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(54) **Process for selective derivatization of taxanes**

Verfahren zur selektiven Derivatisierung von Taxanen

Procédé de dérivation sélective de taxanes

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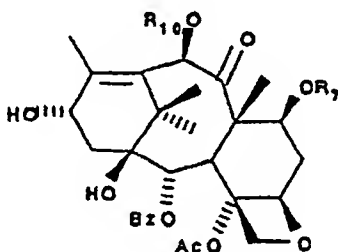
**EP 1 170 293 B1**

## Description

BACKGROUND OF THE INVENTION

[0001] The present invention is directed, in general, to a process for the preparation of taxol and other taxanes, and in particular, to such a process in which the C(7) or C(10) hydroxyl group of a taxane is selectively derivatized.

[0002] 10-DAB (1), which is extracted from the needles of *taxus baccata* L., the English yew, has become a key starting material in the production of taxol and Taxotere, both of which are potent anticancer agents. Conversion of 10-DAB to taxol, Taxotere® and other taxanes having antitumor activity requires protection or derivatization of the C(7) and C(10) hydroxyl groups followed by esterification of the C(13) hydroxyl group to attach an appropriate side chain at that position.



1 R<sub>10</sub> = R<sub>7</sub> = H

2 R<sub>10</sub> = H, R<sub>7</sub> = TES

[0003] Until now, strategies for the preparation of taxol and taxol analogs were based upon the observation of Senilh et al. (*C.R. Acad. Sci. Paris, II*, 1981, 293, 501) that the relative reactivity of the four hydroxyl groups of 10-DAB toward acetic anhydride in pyridine is C(7)-OH > C(10)-OH > C(13)-OH > C(1)-OH. Denis, et. al. reported (*J. Am. Chem. Soc.*, 1988, 110, 5917) selective silylation of the C(7) hydroxyl group of 10-DAB with triethylsilyl chloride in pyridine to give 7-triethylsilyl-10-deacetyl baccatin (III) (2) in 85% yield. Based upon these reports, in those processes in which differentiation of the C(7) and C(10) hydroxyl groups is required (e.g., preparation of taxol from 10-DAB), the C(7) hydroxyl group must be protected (or derivatized) before the C(10) hydroxyl group is protected or derivatized. For example, taxol may be prepared by treating 10-DAB with triethylsilyl chloride to protect the C(7) hydroxyl group, acetylating the C(10) hydroxyl group, attaching the side chain by esterification of the C(13) hydroxyl group, and, finally, removal of protecting groups.

[0004] It is known that taxanes having various substituents bonded to either the C(10) or the C(7) oxygens show anticancer activity. To provide for more efficient synthesis of these materials, it would be useful to have methods which permit more efficient and more highly selective protection or derivatization of the C(10) and the C(7) hydroxyl groups.

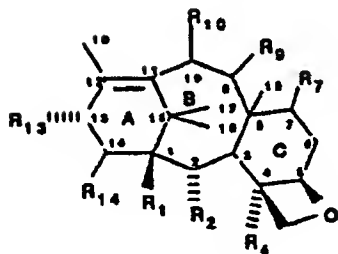
SUMMARY OF THE INVENTION

[0005] Among the objects of the present invention, therefore, is the provision of highly efficient processes for the preparation of taxol and other taxanes through selective derivatization of the C(7) group or the C(10) hydroxyl group of 10-DAB and other taxanes, particularly a process in which the C(10) hydroxyl group is protected or derivatized prior to the C(7) hydroxyl group; and the provision of C(7) or C(10) derivatized taxanes.

[0006] Briefly, therefore, the present invention is directed to a process for the acylation of the C(10) hydroxy group of a taxane. The process comprises forming a reaction mixture containing the taxane and an acylating agent which contains less than one equivalent of a base for each equivalent of taxane, and allowing the taxane to react with the acylating agent to form a C(10) acylated taxane.

[0007] The present invention is further directed to a process for converting the C(7) hydroxy group of a 10-acyloxy-7-hydroxytaxane to an acetal or ketal. The process comprises treating the 10-acyloxy-7-hydroxytaxane with a ketalizing agent in the presence of an acid catalyst to form a C(10) ketalized taxane.

[0008] The process of the present invention preferably produces a taxane having the structure:



wherein

$R_1$  is hydrogen, hydroxy, protected hydroxy, or together with  $R_{14}$  or  $R_2$  forms a carbonate;

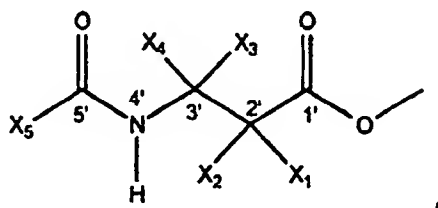
$R_2$  is keto,  $-OT_2$ , acyloxy, or together with  $R_1$  forms a carbonate;

$R_7$  is hydrogen, halogen,  $-OT_7$ ,  $-OCOZ_7$ , or  $-OCOOZ_7$ ;

$R_9$  is hydrogen, keto,  $-OT_9$ ,  $-OCOZ_7$ , or  $-OCOOZ_7$ ;

$R_{10}$  is hydroxy;

$R_{13}$  is hydroxy, protected hydroxy, keto, or



$R_{14}$  is hydrogen,  $-OT_{14}$ , acyloxy, or together with  $R_1$  forms a carbonate;

$T_2$ ,  $T_4$ ,  $T_7$ ,  $T_9$ , and  $T_{14}$  are independently hydrogen or hydroxy protecting group;

$X_1$  is  $-OX_6$ ,  $-SX_7$ , or  $-NX_8X_9$ ;

$X_2$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

$X_3$  and  $X_4$  are independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

$X_5$  is  $-X_{10}$ ,  $-OX_{10}$ ,  $-SX_{10}$ ,  $-NX_8X_{10}$ , or  $-SO_2X_{11}$ ;

$X_6$  is hydrocarbyl, substituted hydrocarbyl, heteroaryl, hydroxy protecting group or a functional group which increases the water solubility of the taxane derivative;

$X_7$  is hydrocarbyl, substituted hydrocarbyl, heteroaryl, or sulfhydryl protecting group;

$X_8$  is hydrogen, hydrocarbyl, or substituted hydrocarbyl;

$X_9$  is an amino protecting group;

$X_{10}$  is hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

$X_{11}$  is hydrocarbyl, substituted hydrocarbyl, heteroaryl,  $-OX_{10}$ , or  $-NX_8X_{14}$ ; and

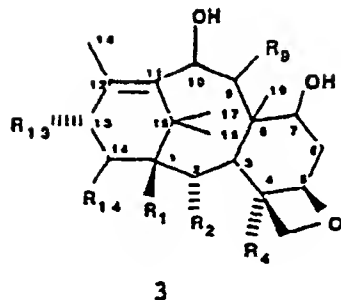
$X_{14}$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

[0009] Other objects and features of this invention will be in part apparent and in part pointed out hereinafter.

#### DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

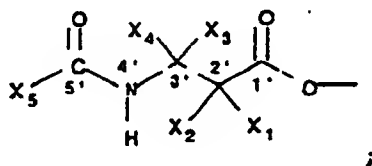
[0010] Among other things, the present invention enables the selective derivatization of the C(10) hydroxyl group of a taxane without first protecting the C(7) hydroxyl group. Stated another way, it has been discovered that the reactivities previously reported for the C(7) and C(10) hydroxyl groups can be reversed, that is, the reactivity of the C(10) hydroxyl group becomes greater than the reactivity of the C(7) hydroxyl group under certain conditions.

[0011] Although the present invention may be used to selectively derivatize a taxane having a hydroxy group at C(7) or C(10), it offers particular advantages in the selective derivatization of taxanes having hydroxy groups at C(7) and C(10), i.e., 7,10-dihydroxy taxanes. In general, 7,10-dihydroxytaxanes which may be selectively derivatized in accordance with the present invention correspond to the following structure:



wherein

- $R_1$  is hydrogen, hydroxy, protected hydroxy, or together with  $R_{14}$  or  $R_2$  forms a carbonate;  
 $R_2$  is keto,  $-OT_2$ , acyloxy, or together with  $R_1$  forms a carbonate;  
 $R_4$  is  $-OT_4$  or acyloxy;  
 $R_9$  is hydrogen, keto,  $-OT_9$ , or acyloxy;  
 $R_{13}$  is hydroxy, protected hydroxy, keto, or

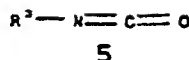
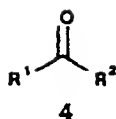


- $R_{14}$  is hydrogen,  $-OT_{14}$ , acyloxy or together with  $R_1$  forms a carbonate;  
 $T_2$ ,  $T_4$ ,  $T_9$ , and  $T_{14}$  are independently hydrogen or hydroxy protecting group;  
 $X_1$  is  $-OX_6$ ,  $-SX_7$ , or  $-NX_8X_9$ ;  
 $X_2$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;  
 $X_3$  and  $X_4$  are independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;  
 $X_5$  is  $X_{10}$ ,  $-OX_{10}$ ,  $-SX_{10}$ ,  $-NX_8X_{10}$ , or  $-SO_2X_{11}$ ;  
 $X_8$  is hydrocarbyl, substituted hydrocarbyl, heteroaryl, hydroxy protecting group or a functional group which increases the water solubility of the taxane derivative;  
 $X_7$  is hydrocarbyl, substituted hydrocarbyl, heteroaryl, or sulfhydryl protecting group;  
 $X_6$  is hydrogen, hydrocarbyl, or substituted hydrocarbyl;  
 $X_9$  is an amino protecting group;  
 $X_{10}$  is hydrocarbyl, substituted hydrocarbyl, or heteroaryl;  
 $X_{11}$  is hydrocarbyl, substituted hydrocarbyl, heteroaryl,  $-OX_{10}$ , or  $-NX_8X_{14}$ ; and  
 $X_{14}$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

#### Selective C(10) Derivatization

[0012] In accordance with the process of the present invention, it has been discovered that the C(10) hydroxyl group of a taxane can be selectively acylated in the absence of a base, preferably in the absence of an amine base. Preferably, therefore, amine bases such as pyridine, triethylamine, dimethylaminopyridine and 2,6-lutidine, if present at all, are present in the reaction mixture in relatively low concentration. Stated another way, if a base is present in the reaction mixture, the molar-ratio of the amine base to the taxane is preferably less than 1:1, more preferably less than 10:1, and most preferably less than 100:1.

[0013] Acylating agents which may be used for the selective acylation of the C(10) hydroxyl group of a taxane include anhydrides, dicarbonates, thiodicarbonates, and isocyanates. In general, the anhydrides, dicarbonates, and thiodicarbonates correspond to structure 4 and the isocyanates correspond to structure 5:



wherein R<sup>1</sup> is -OR<sup>a</sup>, -SR<sup>a</sup>, or R<sup>a</sup>; R<sup>2</sup> is -OC(O)R<sup>b</sup>, -OC(O)OR<sup>b</sup>, -OC(O)SR<sup>b</sup>, -OPOR<sup>b</sup>R<sup>c</sup>, or -OS(O)<sub>2</sub>R<sup>b</sup>; R<sup>3</sup> is hydrocarbyl, substituted hydrocarbyl, or heteroaryl; and R<sup>a</sup>, R<sup>b</sup>, R<sup>c</sup> are independently hydrocarbyl, substituted hydrocarbyl, or heteroaryl. For example, suitable carboxylic acid anhydride acylating agents include acetic anhydride, chloroacetic anhydride, propionic anhydride, benzoic anhydride, and other carboxylic acid anhydrides containing substituted or unsubstituted hydrocarbyl or heteroaryl moieties; suitable dicarbonate acylating reagents include dibenzyl dicarbonate, diallyl dicarbonate, dipropyl dicarbonate, and other dicarbonates containing substituted or unsubstituted hydrocarbyl or heteroaryl moieties; and suitable isocyanate acylating agents include phenyl isocyanate, and other isocyanates containing substituted or unsubstituted hydrocarbyl or heteroaryl moieties. In addition, although the anhydrides, dicarbonates, and thi-

dicarbonates used as acylating agents may be mixed, it is generally preferred that they be symmetrical; that is, R<sup>1</sup> and R<sup>2</sup> are selected such that the molecule is symmetrical (e.g., if R<sup>1</sup> is R<sup>a</sup>, R<sup>2</sup> is -OC(O)R<sup>b</sup> with R<sup>a</sup> being the same as R<sup>b</sup>).

[0014] While the acylation of the C(10) hydroxy group of the taxane will proceed at an adequate rate for many acylating agents, it has been discovered that the reaction rate may be increased by including a Lewis acid in the reaction mixture.

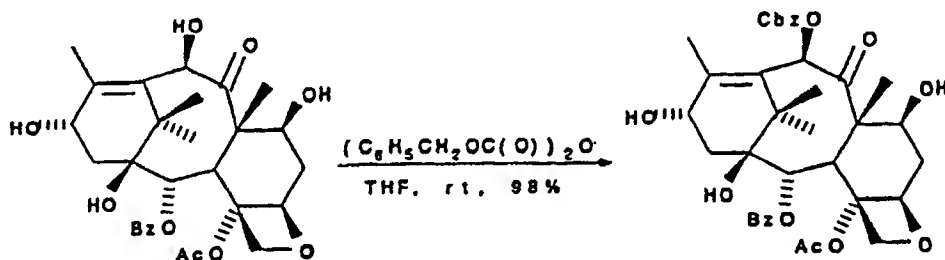
The concentration of the Lewis acid appears not to be narrowly critical; experimental evidence obtained to date suggests it may be present in either a stoichiometric or a catalytic amount. In general, Lewis acids which may be used include triflates and halides of elements of groups IB, IIB, IIIB, IVB, VB, VIB, VIIB, VIII, IIIA, IVA, lanthanides, and actinides of the Periodic Table (American Chemical Society format). Preferred Lewis acids include zinc chloride, stannic chloride, cerium trichloride, cuprous chloride, lanthanum trichloride, dysprosium trichloride, and ytterbium trichloride. Zinc chloride or cerium trichloride is particularly preferred when the acylating agent is an anhydride or dicarbonate. Cuprous chloride is particularly preferred when the acylating agent is an isocyanate.

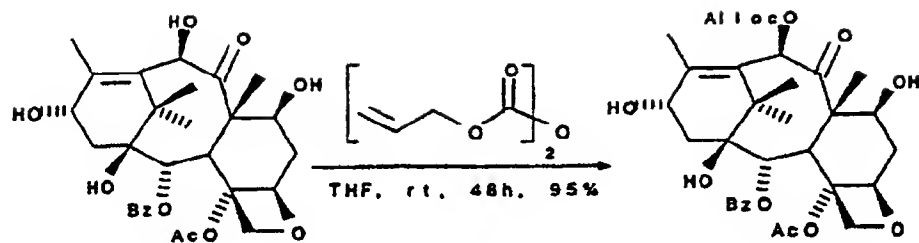
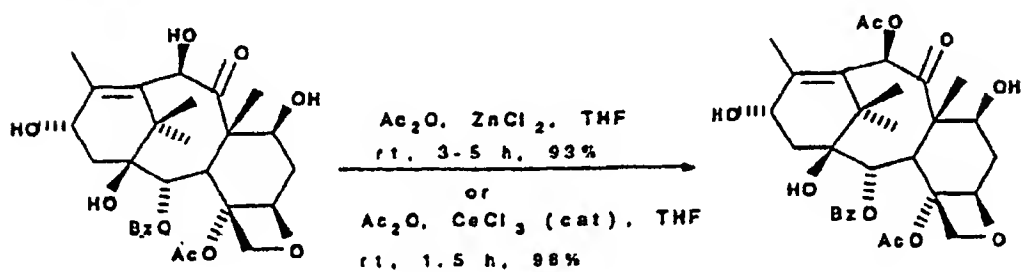
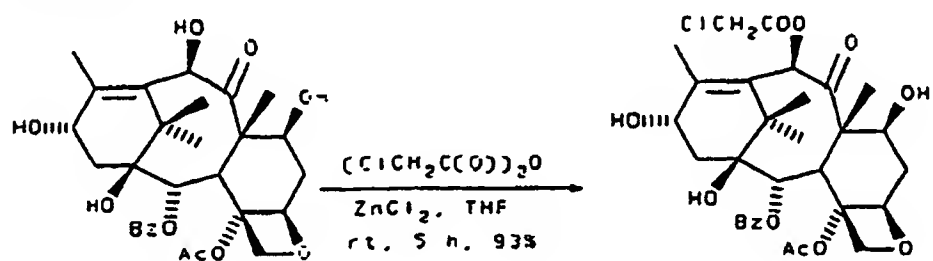
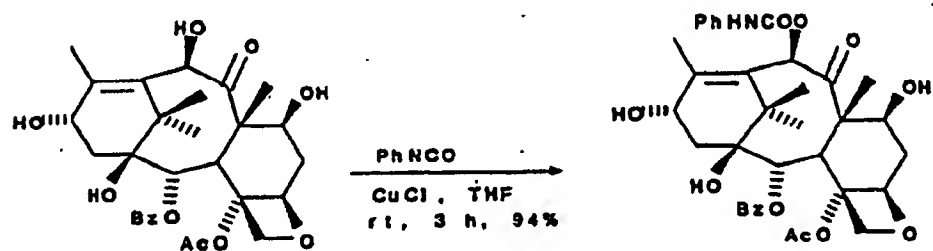
[0015] The solvent for the selective acylation is preferably an ethereal solvent such as tetrahydrofuran. Alternatively, however, other solvents such as ether or dimethoxyethane may be used.

[0016] The temperature at which the C(10) selective acylation is carried out is not narrowly critical. In general, however, it is preferably carried out at room temperature or higher in order for the reaction to proceed at a sufficiently high rate.

[0017] For purposes of illustration, acylating reactions involving dibenzyl dicarbonate, diallyl dicarbonate, acetic anhydride, chloroacetic anhydride and phenyl isocyanate are illustrated in Reaction Schemes 1 through 5 below. In this series of reaction schemes, the taxane which is selectively acylated at the C(10) position is 10-deacetylbaccatin III. It should be understood, however, that these reaction schemes are merely illustrative and that other taxanes having a C(10) hydroxy group, in general, and other 7,10-dihydroxytaxanes, in particular, may be selectively acylated with these and other acylating agents in accordance with the present invention.

Scheme 1



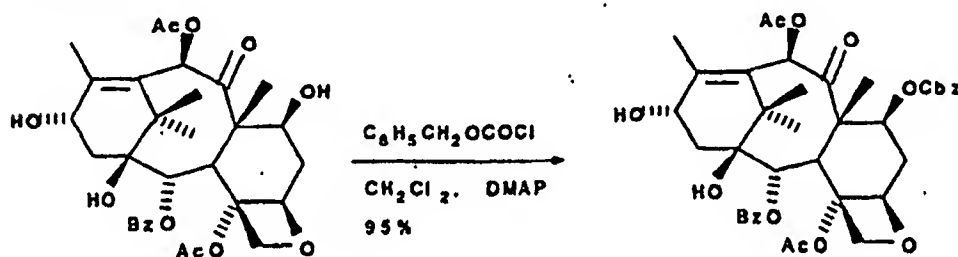
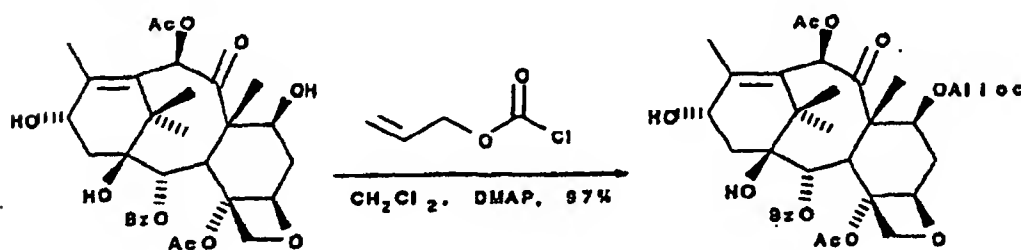
Scheme 2Scheme 3Scheme 4Scheme 5

Selective C(7) Derivatization

[0018] Selective acylation of the C(7) hydroxyl group of a C(10) acylated taxane can be achieved using any of a variety of common acylating agents including, but not limited to, substituted and unsubstituted carboxylic acid derivatives, e.g., carboxylic acid halides, anhydrides, dicarbonates, isocyanates and haloformates. For example, the C(7) hydroxyl group of baccatin III, 10-acyl-10-deacetylbaccatin III or 10-trihydrocarbylsilyl-10-deacetyl baccatin III can be selectively acylated with dibenzyl dicarbonate, diallyl dicarbonate, 2,2,2-trichloroethyl chloroformate, benzyl chloroformate or another common acylating agent.

[0019] In general, acylation of the C(7) hydroxy group of a C(10) acylated taxane are more efficient and more selective than are C(7) acylations of a 7,10-dihydroxy taxane such as 10-DAB, i.e., once the C(10) hydroxyl group has been acylated, there is a significant difference in the reactivity of the remaining C(7), C(13), and C(1) hydroxyl groups (and the C(14) hydroxyl group, if present). These acylation reactions may optionally be carried out in the presence or absence of a base.

[0020] Examples of selective C(7) acylation of a taxane having an acylated C(10) hydroxy group are shown in Reaction Schemes 6 and 7. In these reaction schemes, the taxane which is selectively acylated at the C(7) position is baccatin III. It should be understood, however, that these reaction schemes are merely illustrative and that taxanes having other acyl moieties at C(10) as well as other substituents at other taxane ring positions may be selectively acylated at C(7) with these and other acylating agents in accordance with the present invention.

Scheme 6Scheme 7

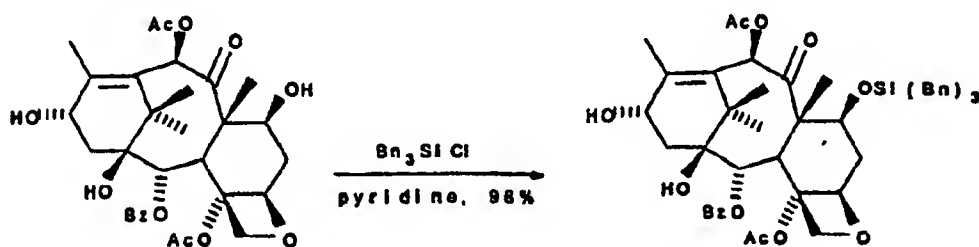
[0021] Alternatively, the C(7) hydroxyl group of a C(10) acylated taxane derivative can be selectively protected using any of a variety of hydroxy protecting groups, such as acetal, ketal, silyl, and removable acyl protecting groups. For example, the C(7) hydroxyl group may be silylated using any of a variety of common silylating agents including, but not limited to, tri(hydrocarbyl)silyl halides and tri(hydrocarbyl)silyl triflates. The hydrocarbyl moieties of these compounds may be substituted or unsubstituted and preferably are substituted or unsubstituted alkyl or aryl. For example, the C(7) hydroxyl group of baccatin III can be selectively silylated using silylating agents such as tribenzylsilyl chloride, trimethylsilyl chloride, triethylsilyl chloride, dimethyl isopropylsilyl chloride, dimethyl phenylsilyl chloride, and the like.

[0022] In general, silylations of the C(7) hydroxy group of a C(10) acylated taxanes are more efficient and more selective than are silylations of a 7,10-dihydroxy taxane such as 10-DAB, i.e., once the C(10) hydroxyl group has been acylated, there is a significant difference in the reactivity of the remaining C(7), C(13), and C(1) hydroxyl groups (and

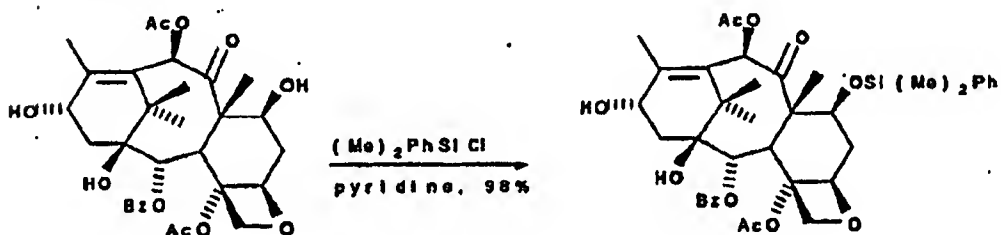
the C(14) hydroxyl group, if present). The C(7) silylation reaction may be carried out under a wide range of conditions, including in the presence or absence of an amine base.

[0023] Examples of selective C(7) silylation of C(10) acylated taxanes are shown in Reaction Schemes 8 through 11. In these reaction schemes, the taxane which is selectively silylated at the C(7) position is baccatin III or another C(10)-acyloxy derivative of 10-deacetylbaccatin III. It should be understood, however, that these reaction schemes are merely illustrative and that other taxanes may be selectively silylated with these and other silylating agents in accordance with the present invention.

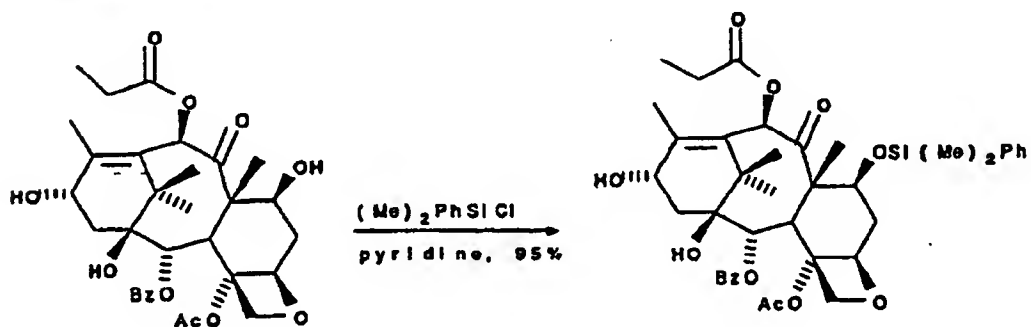
### Scheme 8



### Scheme 9

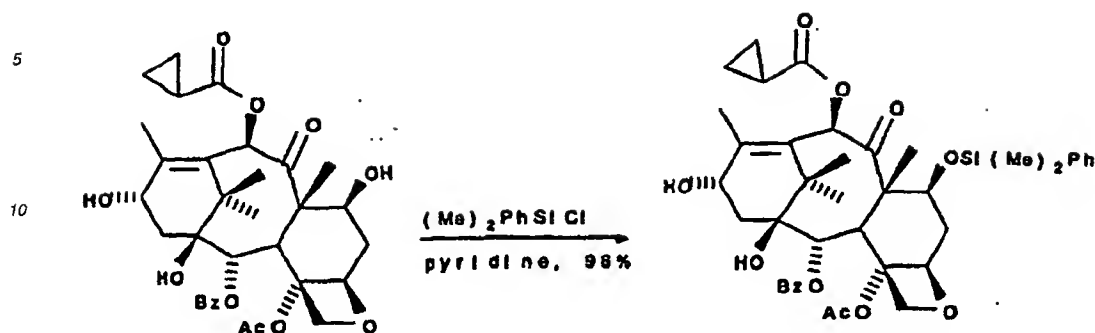


### Scheme 10





Scheme 11

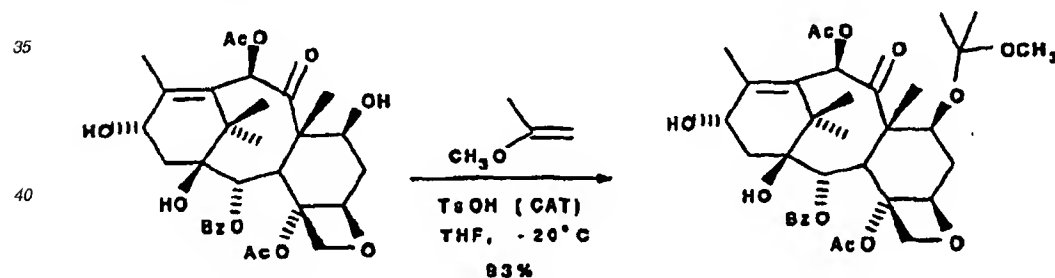


[0024] Alternatively, the C(7) hydroxyl group of C(10) acylated taxanes can be selectively protected using any of a variety of common reagents including, but not limited to, simple acetals, ketals and vinyl ethers, in the presence of an acid catalyst. These reagents (whether acetal, ketal, vinyl ether or otherwise) are referred to herein as "ketalizing agents" and are described in "Protective Groups in Organic Synthesis" by T. W. Greene, John Wiley and Sons, 1981. The acid catalyst used may be an organic or inorganic acid, such as toluenesulfonic acid or camphorsulfonic acid, in at least a catalytic amount. For example, the C(7) hydroxyl group of baccatin III can be selectively ketalized using 2-methoxy propene. Other suitable reagents for the preparation of acetals and ketals include methyl vinyl ether, ethyl vinyl ether, tetrahydropyran, and the like.

[0025] Selective ketalization of the C(7) substituent of a C(10) acylated taxane is more efficient and more selective than it is with 10-DAB, i.e., once the C(10) hydroxyl group has been acylated, there is a large difference in the reactivity of the remaining C(7), C(13), and C(1) hydroxyl groups (and the C(14) hydroxyl group, if present).

[0026] An example of selective formation of a C(7) ketal from baccatin III is illustrated in Reaction Scheme 12. It should be understood, however, that this reaction scheme is merely illustrative and that other taxanes may be selectively ketalized with this and other ketalizing agents in accordance with the present invention.

Scheme 12



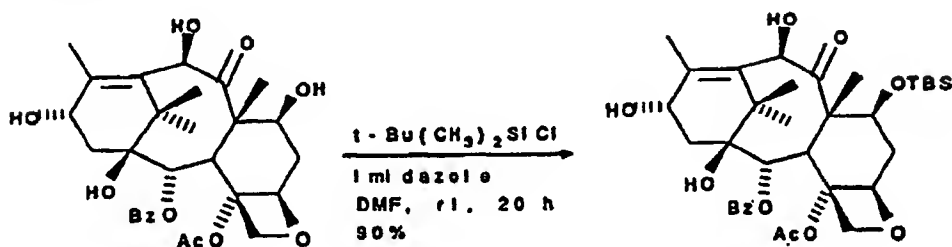
[0027] Under appropriate conditions, the C(7) hydroxyl group of a taxane further comprising a C(10) hydroxyl group can be selectively silylated. Advantageously, these silylations are not limited to silyl groups bearing alkyl substituents having three carbons or less.

[0028] In general, the C(7) hydroxyl group of a taxane can be selectively silylated with a silylating agent which includes the  $-\text{SiR}_j\text{R}_k\text{R}_l$  moiety wherein  $\text{R}_j$ ,  $\text{R}_k$  and  $\text{R}_l$  are independently substituted or unsubstituted hydrocarbyl or heteroaryl, provided that any substituents are other than hydroxyl. In one embodiment of the present invention, if each of  $\text{R}_j$ ,  $\text{R}_k$  and  $\text{R}_l$  is alkyl, then at least one of  $\text{R}_j$ ,  $\text{R}_k$ , and  $\text{R}_l$  comprises a carbon skeleton (i.e., carbon chain or ring(s)) having at least four carbon atoms. Suitable silylating agents include silyl halides and silyl triflates, for example, tri(hydrocarbyl)silyl halides and tri(hydrocarbyl)silyl triflates. The hydrocarbyl substituents of these silylating agents may be substituted or unsubstituted and preferably are substituted or unsubstituted alkyl or aryl.

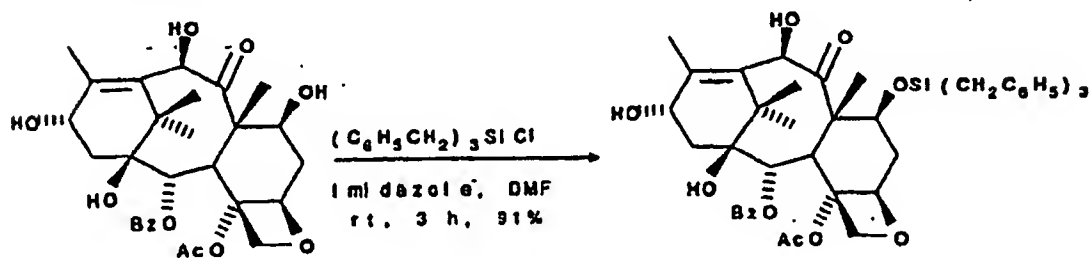
[0029] The selective silylation of the C(7) hydroxy group may be carried out in a solvent, such as dimethyl formamide ("DMF") or pyridine and in the presence of an amine base, such as imidazole or pyridine. Reaction Schemes 13 - 16 illustrate the silylation of the C(7) hydroxy group of 10-DAB in high yield by treating 10-DAB with t-butyltrimethylsilyl chloride, tribenzylsilyl chloride, dimethyl-isopropylsilyl chloride, and dimethylphenylsilyl chloride, respectively. Silylation

under these conditions was surprising in view of the report by Denis, et. al. (*J. Am. Chem. Soc.*, 1988, 110, 5917) that selective formation of 7-TBS-10-DAB was not possible.

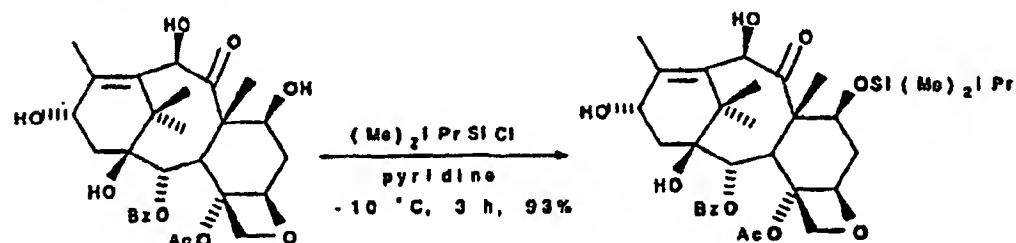
### Scheme 13



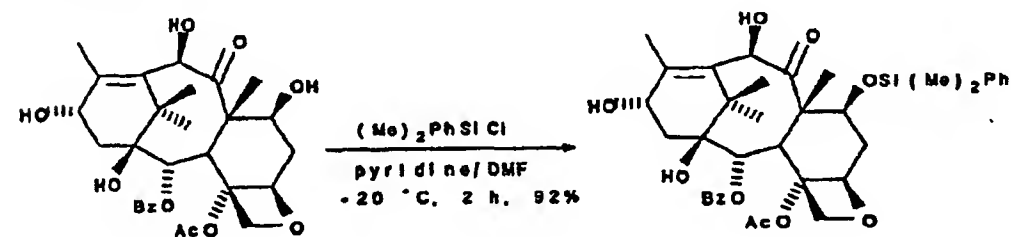
### Scheme 14



### Scheme 15



### Scheme 16



[0030] The process of the present invention can also be used to protect the C(7) hydroxy groups of a 7,10-dihydroxytaxane with different silyl protecting groups. By selecting groups which can be removed under different conditions, the C(7) and C(10) hydroxy groups can be separately accessed for derivatization. These reactions, therefore, increase the

flexibility of the overall process and, enable a higher yield for many of the individual protecting reactions relative to the yield obtained using currently available processes.

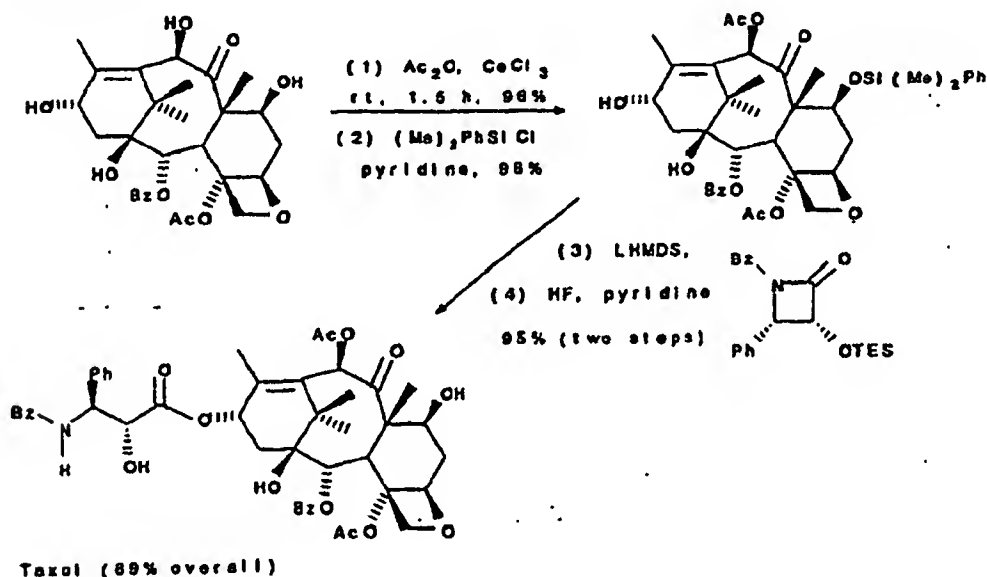
[0031] The methods disclosed herein may be used in connection with a large number of different taxanes obtained from natural or synthetic sources to prepare a wide variety of taxane intermediates which may then be further derivatized.

For example, the methods of the present invention may be effectively used to protect the C(7) and/or C(10) hydroxy functional group prior to the coupling reaction between a C(13) side chain precursor and a taxane to introduce a C(13)  $\beta$ -amido ester side chain, and also prior to the reactions for preparing taxanes having alternative substituents at various locations on the taxane nucleus.

[0032] The attachment of a C(13) side chain precursor to a taxane may be carried out by various known techniques.

For example, a side chain precursor such as an appropriately substituted  $\beta$ -lactam, oxazoline, oxazolidine carboxylic acid, oxazolidine carboxylic acid anhydride, or isoserine derivative may be reacted with a tricyclic or tetracyclic taxane having a C(13) hydroxy, metallic oxide or ammonium oxide substituent to form compounds having a  $\beta$ -amido ester substituent at C(13) as described, for example, Taxol: Science and Applications, M. Suffness, editor, CRC Press (Boca Roton, FL) 1995, Chapter V, pages 97-121. For example, the synthesis of taxol from 10-DAB is illustrated in Reaction Scheme 17. It should be noted that while a  $\beta$ -lactam and 10-DAB are used in this reaction scheme, other side chain precursors and other taxanes could be substituted therefor without departing from the present invention.

**Scheme 17**



[0033] The process illustrated in Reaction Scheme 17 is significantly more efficient than any other currently known process, due to the high yields and selectivity of the cerium trichloride catalyzed acetylation of the C(10) hydroxyl group of 10-DAB and the subsequent silylation of the C(7) hydroxyl group. The synthesis proceeds in four steps and 89% overall yield.

[0034] The protected taxane derivatives or the intermediates or starting materials used in the preparation of such protected taxane derivatives can be further modified to provide for alternative substituents at various positions of the taxane.

[0035] Taxanes having C(2) and/or C(4) substituents other than benzoyloxy and acetoxy, respectively, can be prepared from baccatin III, 10-DAB and other taxanes as more fully described in PCT Patent Application WO 94/01223. In general, the C(2) and C(4) acyloxy substituents are treated with lithium aluminum hydride or another suitable reducing agent to form hydroxy groups at C(2) and C(4) which may then be reacted, for example, with carboxylic acid halides (optionally after protection of the C(2) hydroxy group together with the C(1) hydroxy group with a 1,2-carbonate protecting group) to obtain the desired C(2) and C(4) derivatives.

[0036] Taxanes having C(7) substituents other than hydroxy and acyloxy as described herein can be prepared from baccatin III, 10-DAB, and other taxanes as more fully described in PCT Patent Application WO 94/17050. For example,

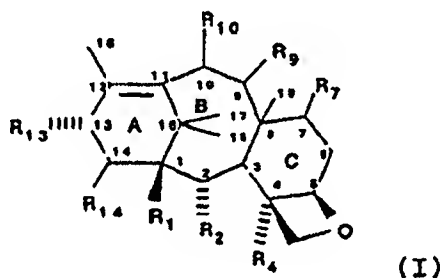
a C(7) xanthate may be subjected to tin hydride reduction to yield the corresponding C(7) dihydro taxane. Alternatively, C(7) fluoro-substituted taxanes can be prepared by treatment of C(13)-triethylsilyl-protected baccatin III with 2-chloro-1,1,2-trifluoroethylamine at room temperature in THF solution. Other baccatin derivatives with a free C(7) hydroxyl group behave similarly. Alternatively, 7-chloro baccatin III can be prepared by treatment of baccatin III with methanesulfonyl chloride and triethylamine in methylene chloride solution containing an excess of triethylamine hydrochloride.

**[0037]** Taxanes having C(9) substituents other than keto can be prepared from baccatin III, 10-DAB and other taxanes as more fully described in PCT Patent Application WO 94/20088. In general, the C(9) keto substituent of the taxane is selectively reduced to yield the corresponding C(9)  $\beta$ -hydroxy derivative with a borohydride, preferably tetrabutylammonium borohydride ( $\text{Bu}_4\text{NBH}_4$ ) or triacetoxyborohydride. The C(9)  $\beta$ -hydroxy derivative can then be protected at C(7) with a hydroxy protecting group and the C(9) hydroxy group can be acylated following the methods described herein for acylation of the C(7) hydroxy group. Alternatively, reaction of 7-protected-9 $\beta$ -hydroxy derivative with KH causes the acetate group (or other acyloxy group) to migrate from C(10) to C(9) and the hydroxy group to migrate from C(9) to C(10), thereby yielding a 10-desacetyl derivative, which can be acylated as described elsewhere herein.

**[0038]** Taxanes having C(10) substituents other than hydroxy, acyloxy or protected hydroxy as described herein may be prepared as more fully described in PCT Patent Application WO 94/15599 and other literature references. For example, taxanes having a C(10) keto substituent can be prepared by oxidation of 10-desacetyl taxanes. Taxanes which are dihydro substituted at C(10) can be prepared by reacting a C(10) hydroxy or acyloxy substituted taxane with samarium diiodide.

**[0039]** Taxanes having a C(14) substituent other than hydrogen may also be prepared. The starting material for these compounds may be, for example, a hydroxylated taxane (14-hydroxy-10-deacetyl baccatin III) which has been discovered in an extract of yew needles (C&EN, p 36-37, April 12, 1993). Derivatives of this hydroxylated taxane having the various C(2), C(4), C(7), C(9), C(10), C3' and C5' functional groups described above may also be prepared by using this hydroxylated taxane. In addition, the C(14) hydroxy group together with the C(1) hydroxy group of 10-DAB can be converted to a 1,2-carbonate as described in C&EN or it may be converted to a variety of esters or other functional groups as otherwise described herein in connection with the C(2), C(4), C(9) and C(10) substituents.

**[0040]** The process of the present invention thus enables the preparation of taxanes having the following structure:



wherein

M comprises ammonium or is a metal;

R<sub>1</sub> is hydrogen, hydroxy, protected hydroxy, or together with R<sub>14</sub> or R<sub>2</sub> forms a carbonate;

R<sub>2</sub> is keto, -OT<sub>2</sub>, acyloxy, or together with R<sub>1</sub> forms a carbonate;

R<sub>4</sub> is -OT<sub>4</sub> or acyloxy;

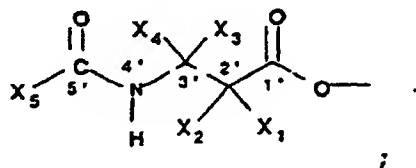
R<sub>7</sub> is hydrogen, halogen, -OT<sub>7</sub>, or acyloxy;

R<sub>9</sub> is hydrogen, keto, -OT<sub>9</sub>, or acyloxy;

R<sub>10</sub> is hydrogen, keto, -OT<sub>10</sub>, or acyloxy;

R<sub>7</sub>, R<sub>9</sub>, and R<sub>10</sub> independently have the alpha or beta stereochemical configuration;

R<sub>13</sub> is hydroxy, protected hydroxy, keto, MO- or



$R_{14}$  is hydrogen,  $-OT_{14}$ , acyloxy, or together with  $R_1$  forms a carbonate;

$T_2, T_4, T_7, T_9, T_{10}$ , and  $T_{14}$  are independently hydrogen or hydroxy protecting group;

$X_1$  is  $-OX_6$ ,  $-SX_7$ , or  $-NX_8X_9$ ;

$X_2$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

5  $X_3$  and  $X_4$  are independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

$X_5$  is  $-X_{10}$ ,  $-OX_{10}$ ,  $-SX_{10}$ ,  $-NX_8X_{10}$ , or  $-SO_2X_{11}$ ;

$X_6$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, heteroaryl, hydroxy protecting group, or a functional group which increases the water solubility of the taxane derivative;

$X_7$  is hydrocarbyl, substituted hydrocarbyl, heteroaryl, or sulfhydryl protecting group;

10  $X_8$  is hydrogen, hydrocarbyl, or substituted hydrocarbyl;

$X_9$  is an amino protecting group;

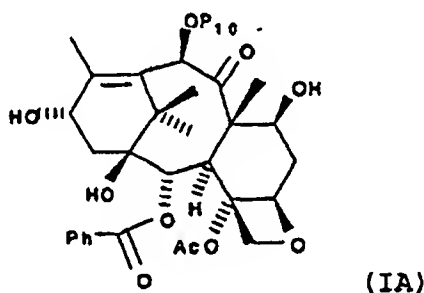
$X_{10}$  is hydrocarbyl, substituted hydrocarbyl, or heteroaryl;

$X_{11}$  is hydrocarbyl, substituted hydrocarbyl, heteroaryl,  $-OX_{10}$ , or  $-NX_8X_{14}$ ;

$X_{14}$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

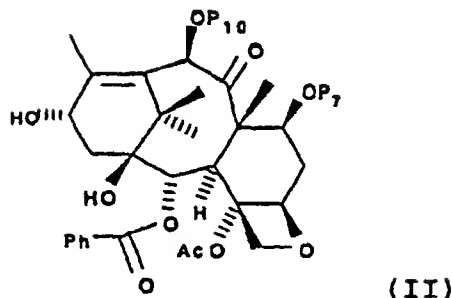
15 **[0041]** In one embodiment of the present invention, the substituents of the taxane (other than the C(7), C(10) and C(13) substituents) correspond to the substituents present on baccatin III or 10-DAB. That is,  $R_{14}$  is hydrogen,  $R_9$  is keto,  $R_4$  is acetoxy,  $R_2$  is benzoyloxy, and  $R_1$  is hydroxy. In other embodiments, the taxane has a structure which differs from that of taxol or Taxotere® with respect to the C(13) side chain and at least one other substituent. For example,  $R_{14}$  may be hydroxy;  $R_2$  may be hydroxy,  $-OCOZ_2$  or  $-OCOOZ_{22}$  wherein  $Z_2$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl and  $Z_{22}$  is hydrocarbyl, substituted hydrocarbyl, or heteroaryl;  $R_4$  may be hydroxy,  $-OCOZ_4$  or  $-OCOOZ_{44}$  wherein  $Z_4$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl and  $Z_{44}$  is hydrocarbyl, substituted hydrocarbyl, or heteroaryl;  $R_7$  may be hydrogen, hydroxy,  $-OCOZ_7$  or  $-OCOOZ_{77}$  wherein  $Z_7$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl and  $Z_{77}$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;  $R_9$  may be hydrogen, hydroxy,  $-OCOZ_9$  or  $-OCOOZ_{99}$  wherein  $Z_9$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl and  $Z_{99}$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl, and  $R_{10}$  may be hydrogen, hydroxy,  $-OCOZ_{10}$  or  $-OCOOZ_{1010}$  wherein  $Z_{10}$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl and  $Z_{1010}$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

**[0042]** In a preferred embodiment, the taxane has the formula



wherein  $P_{10}$  is acyl, said acyl comprising at least three carbon atoms or two carbon atoms and a nitrogen, oxygen or sulfur atom. Stated another way,  $-OP_{10}$  is other than acetoxy. More preferably,  $P_{10}$  is  $-(C=O)R_A$ ,  $-(C=O)OR_B$ , or  $-(C=O)NR_C$  wherein  $R_A$  is substituted or unsubstituted hydrocarbyl or heteroaryl, said unsubstituted hydrocarbyl comprising at least two carbon atoms;  $R_B$  and  $R_C$  are independently substituted or unsubstituted hydrocarbyl. Still more preferably,  $R_A$  is substituted or unsubstituted alkyl or aryl, said unsubstituted alkyl comprising at least two carbon atoms; and  $R_B$  and  $R_C$  are independently substituted or unsubstituted alkyl or aryl.

**[0043]** In another embodiment of the invention, the taxane has the formula



wherein  $P_7$  and  $P_{10}$  are independently substituted or unsubstituted acyl. In this embodiment,  $P_7$  and  $P_{10}$  are preferably different.

#### Definitions

[0044] As used herein, the terms "selective" and "selective derivatization" shall mean that the desired product is preferentially formed over any other by-product. Preferably, the desired product is present in a molar ratio of at least 9:1 relative to any other by-product and, more preferably, is present in a molar ratio of at least 20:1 relative to any other by-product.

[0045] In addition, "Ph" means phenyl; "Bz" means benzoyl; "Bn" means benzyl; "Me" means methyl; "Et" means ethyl; "iPr" means isopropyl; "tBu" and "t-Bu" means tert-butyl; "Ac" means acetyl; "TES" means triethylsilyl; "TMS" means trimethylsilyl; "TBS" means  $\text{Me}_2\text{t-BuSi-}$ ; "CDI" means carbonyl diimidazole; "BOM" means benzyloxymethyl; "DBU" means diazabicycloundecane; "DMAP" means p-dimethylamino pyridine; "LHMDS" or "LiHMDS" means lithium hexamethyldisilazide; "DMF" means dimethylformamide; "10-DAB" means 10-desacetylbaccatin III; "Cbz" means benzyloxycarbonyl; "Alloc" means allyloxycarbonyl; "THF" means tetrahydrofuran; "BOC" means benzyloxycarbonyl; "PNB" means para-nitrobenzyl; "Troc" means 2,2,2-trichloroethoxycarbonyl; "EtOAc" means ethyl acetate; "THF" means tetrahydrofuran; "protected hydroxyl" means -OP wherein P is a hydroxyl protecting group; and "hydroxyl protecting group" includes, but is not limited to, acetals having two to ten carbons, ketals having two to ten carbons, and ethers, such as methyl, t-butyl, benzyl, p-methoxybenzyl, p-nitrobenzyl, allyl, trityl, methoxymethyl, methoxyethoxymethyl, ethoxyethyl, methoxypropyl, tetrahydropyranyl, tetrahydrothiopyranyl; and trialkylsilyl ethers such as trimethylsilyl ether, triethylsilyl ether, dimethylarylsilyl ether, trisopropylsilyl ether and t-butyl dimethylsilyl ether; esters such as benzoyl, acetyl, phenylacetyl, formyl, mono-, di-, and trihaloacetyl such as chloroacetyl, dichloroacetyl, trichloroacetyl, trifluoroacetyl; and carbonates including but not limited to alkyl carbonates having from one to six carbon atoms such as methyl, ethyl, n-propyl, isopropyl, n-butyl, t-butyl; isobutyl, and n-pentyl; alkyl carbonates having from one to six carbon atoms and substituted with one or more halogen atoms such as 2,2,2-trichloroethoxymethyl and 2,2,2-trichloroethyl; alkenyl carbonates having from two to six carbon atoms such as vinyl and allyl; cycloalkyl carbonates having from three to six carbon atoms such as cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl; and phenyl or benzyl carbonates optionally substituted on the ring with one or more  $\text{C}_{1-6}$  alkoxy, or nitro. Other hydroxyl protecting groups may be found in "Protective Groups in Organic Synthesis" by T. W. Greene, John Wiley and Sons, 1981, and Second Edition, 1991.

[0046] The "hydrocarbon" and "hydrocarbyl" moieties described herein are organic compounds or radicals consisting exclusively of the elements carbon and hydrogen. These moieties include alkyl, alkenyl, alkynyl, and aryl moieties. These moieties also include alkyl, alkenyl, alkynyl, and aryl moieties substituted with other aliphatic or cyclic hydrocarbyl groups, and include alkaryl, alkenaryl and alkynaryl. Preferably, these moieties comprise 1 to 20 carbon atoms.

[0047] The alkyl groups described herein are preferably lower alkyl containing from one to six carbon atoms in the principal chain and up to 20 carbon atoms. They may be straight, branched chain or cyclic and include methyl, ethyl, propyl, isopropyl, butyl, hexyl and the like. They may be substituted with aliphatic or cyclic hydrocarbyl radicals.

[0048] The alkenyl groups described herein are preferably lower alkenyl containing from two to six carbon atoms in the principal chain and up to 20 carbon atoms. They may be straight or branched chain and include ethenyl, propenyl, isopropenyl, butenyl, isobutenyl, hexenyl, and the like. They may be substituted with aliphatic or cyclic hydrocarbyl radicals.

[0049] The alkynyl groups described herein are preferably lower alkynyl containing from two to six carbon atoms in the principal chain and up to 20 carbon atoms. They may be straight or branched chain and include ethynyl, propynyl, butynyl, isobutynyl, hexynyl, and the like. They may be substituted with aliphatic or cyclic hydrocarbyl radicals.

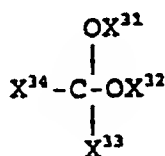
[0050] The aryl moieties described herein contain from 6 to 20 carbon atoms and include phenyl. They may be hydrocarbyl substituted with the various substituents defined herein. Phenyl is the more preferred aryl.

[0051] The heteroaryl moieties described are heterocyclic compounds or radicals which are analogous to aromatic compounds or radicals and which contain a total of 5 to 20 atoms, usually 5 or 6 ring atoms, and at least one atom other than carbon, such as furyl, thienyl, pyridyl and the like. The heteroaryl moieties may be substituted with hydrocarbyl, heterosubstituted hydrocarbyl or hetero-atom containing substituents with the hetero-atoms being selected from the group consisting of nitrogen, oxygen, silicon, phosphorous, boron, sulfur, and halogens. These substituents include hydroxy; lower alkoxy such as methoxy, ethoxy, butoxy; halogen such as chloro or fluoro; ethers; acetals; ketals; esters; heteroaryl such as furyl or thienyl; alkanoxy; acyl; acyloxy; nitro; amino; and amido.

[0052] The substituted hydrocarbyl moieties described herein are hydrocarbyl moieties which are substituted with at least one atom other than carbon and hydrogen, including moieties in which a carbon chain atom is substituted with a hetero atom such as nitrogen, oxygen, silicon, phosphorous, boron, sulfur, or a halogen atom. These substituents include hydroxy; lower alkoxy such as methoxy, ethoxy, butoxy; halogen such as chloro or fluoro; ethers; acetals; ketals; esters; heteroaryl such as furyl or thienyl; alkanoxy; acyl; acyloxy; nitro; amino; and amido.

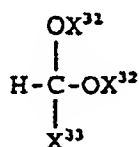
[0053] The acyl moieties and the acyloxy moieties described herein contain hydrocarbyl, substituted hydrocarbyl or heteroaryl moieties. In general, they have the formulas  $-C(O)G$  and  $-OC(O)G$ , respectively, wherein G is substituted or unsubstituted hydrocarbyl, hydrocarbyloxy, hydrocarbylamino, hydrocarbylthio or heteroaryl.

[0054] The ketal moieties described herein have the general formula



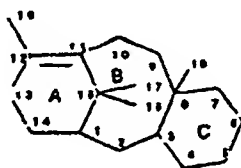
wherein  $X^{31}$ ,  $X^{32}$ ,  $X^{33}$  and  $X^{34}$  are independently hydrocarbyl, substituted hydrocarbyl or heteroaryl moieties. They may be optionally substituted with the various substituents defined herein. The ketal moieties are preferably substituted or unsubstituted alkyl or alkenyl, and more preferably substituted or unsubstituted lower ( $C_1$ - $C_6$ ) alkyl. These ketal moieties additionally may encompass sugars or substituted sugars and include ketal moieties prepared from sugars or substituted sugars such as glucose and xylose. When a ketal moiety is incorporated into a taxane of the present invention as a C(7) hydroxy protecting group, then either  $X^{31}$  or  $X^{32}$  represents the taxane moiety.

[0055] The acetal moieties described herein have the general formula



wherein  $X^{31}$ ,  $X^{32}$  and  $X^{33}$  are independently hydrocarbyl, substituted hydrocarbyl or heteroaryl moieties. They may be optionally substituted with the various substituents defined herein other than hydroxyl. The acetal moieties are preferably substituted or unsubstituted alkyl or alkenyl, and more preferably substituted or unsubstituted lower ( $C_1$ - $C_6$ ) alkyl. These acetal moieties additionally may encompass sugars or substituted sugars and include acetal moieties prepared from sugars or substituted sugars such as glucose and xylose. When an acetal moiety is incorporated into a taxane of the present invention as a C(7) hydroxy protecting group, then either  $X^{31}$  or  $X^{32}$  represents the taxane moiety.

[0056] The term "taxane" as used herein, denotes compounds containing the A, B and C rings (with numbering of the ring positions shown herein):



[0057] The following examples illustrate the invention.

## Example 1

## A. Selective acylation of a C(10) hydroxyl group:

5 **[0058]** *10-Cbz-10-DAB*. To a solution of 10-DAB (30 mg, 0.055 mmol) in THF (1 mL) at room temperature was added dibenzyl pyrocarbonate (320 mg, 1.1 mmol, 20 equiv) under N<sub>2</sub>. The reaction mixture was stirred at room temperature for 24 h. EtOAc (10 mL) was added, and the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:1) as the eluent and dried *in vacuo* overnight to give

10 10-cbz-10-DAB as a colorless solid: yield, 37 mg (98%). mp 205-206 °C;  $[\alpha]_{\text{D}}^{25}$  -63° (CHCl<sub>3</sub>, c = 0.41); <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 1.11 (s, 3 H, Me17), 1.13(s, 3 H, Me16), 1.58(s, 1 H, 1-OH), 1.71 (s, 3 H, Me19), 1.89(ddd, J = 14.7, 10.9, 2.3 Hz, 1 H, H6b), 2.00(d, J = 5.1 Hz, 1 H, 13-OH), 2.08(d, J = 1.0, 3 H, Me18), 2.28(s, 3 H, 4-Ac), 2.30(m, 2 H, H14a, H14b), 2.43(d, J = 4.1 Hz, 1 H, 7-OH), 2.58(ddd, J = 14.7, 9.6, 6.6 Hz, 1 H, H6a), 3.88(d, J = 6.9 Hz, 1 H, H3), 4.19(d, J = 8.6 Hz, 1 H, H20b), 4.31(d, J = 8.6 Hz, 1 H, H20a), 4.44(ddd, J = 10.9, 6.6, 4.1 Hz, 1 H, H7), 4.89(m, 1 H, H13), 4.98(dd, J = 9.6, 2.3 Hz, 1 H, H5), 5.23(d, J = 12.1, 1 H, CHH'OC(O)), 5.26(d, J = 12.1, 1 H, CHH'OC(O)), 5.65

15 (d, J = 6.9 Hz, 1 H, H2), 6.19(s, 1 H, H10), 7.35-7.44(m, 5 H, PhCH<sub>2</sub>O), 7.48(dd, J = 8.1, 7.6 Hz, 2 H, benzoate, *m*), 7.60(tt, J = 7.6, 1.0 Hz, 1 H, benzoate, *p*), 8.11(d, J = 8.1, 1.0 Hz, 2 H, benzoate, *o*) ppm. <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>) δ 9.1 (Me (19)), 15.3 (Me (18)), 20.7(4-Ac), 22.3, 26.7(Me16, Me17), 35.5(C(6)), 38.6(C(14)), 42.5(C(15)), 46.1(C(3)), 58.7(C(8)), 67.9(C(13)), 70.5(OCH<sub>2</sub>Ph), 72.2, 75.0, 76.5(C(7), C(2), C(20)), 79.0, 79.1 (C(1), C(10)), 80.9(C(4)), 84.5 (C(5)), 128.6, 128.8, 129.7, 130.3, 131.9, 133.8(OCH<sub>2</sub>Ph, benzoate), 135.1(C(11)), 147.5(C(12)), 155.6(OC(O)O), 167.4 (benzoate), 171.0(4-Ac), 204.7(C(9))ppm. Anal. Calcd for C<sub>37</sub>H<sub>42</sub>O<sub>12</sub>·1/2H<sub>2</sub>O: C, 64.62; H, 6.30. Found: C, 64.34; H, 6.31.

20 **[0059]** *10-Alloc-10-DAB*. To a solution of 10-DAB (30 mg, 0.055 mmol) in THF (1 mL) at room temperature was added diallyl pyrocarbonate (366 mL, 2.2 mmol, 40 equiv) under N<sub>2</sub>. The reaction mixture was stirred at room temperature for 48 h. TLC analysis indicated the presence of the desired product along with unreacted starting material. EtOAc (20 mL) was added, and the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated. The residue was purified by flash column chromatography using EtOAc: hexanes (1:1) as the eluent and dried *in vacuo* overnight to give 10-alloc-10-DAB as a colorless solid: yield, 23 mg (67%, 95% at the conversion of 70%). The recovered 10-DAB, 9 mg (30%). 10-alloc-10-DAB: mp 201-203 °C;  $[\alpha]_{\text{D}}^{25}$  -81° (CHCl<sub>3</sub>, c = 0.53); <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 1.11 (s, 3 H, Me17), 1.12(s, 3 H, Me16), 1.60(s, 1 H, 1-OH), 1.69(s, 3 H, Me19), 1.87(ddd, J = 14.7, 11.0, 2.1 Hz, 1 H, H6b), 2.05(d, J = 5.1 Hz, 1 H, 13-OH), 2.08(d, J = 1.2, 3 H, Me18), 2.28(s, 3 H, 4-Ac), 2.29(m, 2 H, H14a, H14b), 2.47(d, J = 4.2 Hz, 1 H, 7-OH), 2.57(ddd, J = 14.7, 9.6, 6.7 Hz, 1 H, H6a), 3.86(d, J = 7.0 Hz, 1 H, H3), 4.16(d, J = 8.4 Hz, 1 H, H20b), 4.31(d, J = 8.4 Hz, 1 H, H20a), 4.44(ddd, J = 11.0, 6.7, 4.2 Hz, 1 H, H7), 4.70(br d, J = 5.9 Hz, 2 H, CHH' = CHCH<sub>2</sub>O), 4.90(m, 1H, H13), 4.97(dd, J = 9.6, 2.1 Hz, 1 H, H5), 5.32(dd, J = 10.4, 1.2 Hz, 1 H, CHH' = CHCH<sub>2</sub>O), 5.42(dd, J = 17.2, 1.2 Hz, 1 H, CHH' = CHCH<sub>2</sub>O), 5.63(d, J = 7.0 Hz, 1

35 H, H2), 5.98(ddt, J = 17.2, 10.4, 5.9 Hz, 1 H, CHH' = CHCH<sub>2</sub>O), 6.16(s, 1 H, H10), 7.48(dd, J = 8.1, 7.5 Hz, 2 H, benzoate, *m*), 7.60(tt, J = 7.5, 1.2 Hz, 1 H, benzoate, *p*), 8.11(d, J = 8.1, 1.2 Hz, 2 H, benzoate, *o*) ppm; <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>) δ 9.1(Me(19)), 15.3(Me(18)), 20.7(4-Ac), 22.3, 26.7(Me16, Me17), 35.5(C(6)), 38.6(C(14)), 42.5(C(15)), 46.1(C(3)), 58.7 (C(8)), 67.9(C(13)), 69.3(CH<sub>2</sub>=CHCH<sub>2</sub>O), 72.1, 75.0, 76.5(C(7), C(2), C(20)), 79.0, 79.1(C(1), C(10)), 80.9(C(4)), 84.5 (C(5)), 119.6 (CH<sub>2</sub>=CHCH<sub>2</sub>O), 128.8, 129.7, 130.3, 133.8 (benzoate), 131.4, 131.9 (CH<sub>2</sub>=CHCH<sub>2</sub>O, C(11)), 147.5(C(12)), 155.4(OC(O)O), 167.4(benzoate), 170.9(4-Ac), 204.7(C(9))ppm. Anal. Calcd for C<sub>33</sub>H<sub>40</sub>O<sub>12</sub>: C, 63.05; H, 6.41. Found: C, 62.77; H, 6.48.

B. Selective acylation of a C(10) hydroxyl group using ZnCl<sub>2</sub>:

45 **[0060]** *baccatin III*. To a solution of 10-DAB (100 mg, 0.184 mmol) in THF (6 mL) at room temperature was added a mixture of acetic anhydride (6.5 mL) and ZnCl<sub>2</sub>/ THF solution (0.5 M, 726 mL, 0.368 mmol, 2 equiv) under N<sub>2</sub>. The reaction mixture was stirred at room temperature for 4 h. Then the reaction mixture was diluted with EtOAc (100 mL), washed with saturated aqueous NaHCO<sub>3</sub> solution (40 mLx3), brine. The organic layer was dried over Na<sub>2</sub>SO<sub>4</sub>, concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:1)

50 as the eluent and dried *in vacuo* to give baccatin III as a colorless solid: yield, 100 mg (93%). mp 237-238 °C dec (ref 236-238 °C dec);  $[\alpha]_{\text{D}}^{25}$  -63° (CH<sub>3</sub>OH, c = 0.45) (ref  $[\alpha]_{\text{D}}^{25}$  -54°, CH<sub>3</sub>OH); <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 1.11(s, 6 H, Me16, Me17), 1.61(s, 1 H, 1-OH), 1.67(s, 3 H, Me19), 1.87(ddd, J = 14.7, 10.9, 2.1 Hz, 1 H, H6b), 2.05(d, J = 3.8 Hz, 1 H, 13-OH), 2.05(s, 3 H, Me18), 2.24(s, 3 H, 10-Ac), 2.28(s, 3 H, 4-Ac), 2.30(m, 2 H, H14a, H14b), 2.47(d, J = 4.2 Hz, 1 H, 7-OH), 2.57(ddd, J = 14.7, 9.4, 6.7 Hz, 1 H, H6a), 3.89(d, J = 7.0 Hz, 1 H, H3), 4.16(d, J = 8.4 Hz, 1 H, H20b), 4.31(d, J = 8.4 Hz, 1 H, H20a), 4.47(ddd, J = 10.9, 6.7, 4.2 Hz, 1 H, H7), 4.90(m, 1 H, H13), 4.99(dd, J = 9.4, 2.1 Hz, 1 H, H5), 5.63(d, J = 7.0 Hz, 1 H, H2), 6.33(s, 1 H, H10), 7.48(dd, J = 7.8, 7.8 Hz, 2 H, benzoate, *m*), 7.61(dd, J = 7.8, 7.4 Hz, 1 H, benzoate, *p*), 8.11(d, J = 7.4 Hz, 2 H, benzoate, *o*)ppm. <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>) δ 9.4(Me(19)), 15.6(Me(18)), 20.9(4-Ac, 10-Ac), 22.6, 27.0(Me16, Me17), 35.6(C(6)), 38.6(C(14)), 42.7(C(15)), 46.1(C(3)), 58.8(C(8)), 68.0(C(13)),

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72.3, 75.0, 76.2, 76.4(C(7), C(2), C(10), C(20)), 79.1(C(1)), 80.9(C(4)), 84.5(C(5)), 128.6, 129.4, 130.1, 133.7(benzoate), 132.0(C(11)), 146.3(C(12)), 167.1(benzoate), 170.7, 171.3(10-Ac, 4-Ac), 204.1(C(9))ppm.

**[0061] 10-Chloroacetyl-10-DAB.** To a solution of 10-DAB (116 mg, 0.21 mmol) in THF (3 mL) at room temperature was added a mixture of chloroacetic anhydride (2.8 g, 16.3 mmol, 78 equiv) and  $\text{ZnCl}_2/\text{THF}$  solution (0.5 M, 0.85 mL, 0.42 mmol, 2 equiv) via a syringe under  $\text{N}_2$ . The reaction mixture was stirred at room temperature for 5 h. The reaction mixture was poured into a mixture of EtOAc (200 mL) and saturated aqueous  $\text{NaHCO}_3$  solution (100 mL). The organic layer was separated, and the aqueous layer was extracted with EtOAc (100 mLx3). The organic solution was combined, dried over  $\text{Na}_2\text{SO}_4$ , filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:1) as the eluent and dried overnight in *vacuo* to give 10-chloro-acetyl-10-DAB as a colorless solid: yield, 123 mg (93%). mp 231-233 °C dec;  $[\alpha]_{\text{D}}^{25} -66^\circ$  (EtOAc,  $c = 0.45$ );  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  1.11 (s, 3 H, Me17), 1.12(s, 3 H, Me16), 1.63 (s, 1 H, 1-OH), 1.69 (s, 3 H, Me19), 1.89(ddd,  $J = 14.6, 10.9, 2.1$  Hz, 1 H, H6b), 2.07(d,  $J = 5.2$  Hz, 1 H, 13-OH), 2.09(d,  $J = 1.2, 3$  Hz, Me18), 2.12(d,  $J = 4.5$  Hz, 1 H, 7-OH), 2.29(s, 3 H, 4-Ac), 2.30(m, 2 H, H14a, H14b), 2.58(ddd,  $J = 14.6, 9.7, 6.7$  Hz, 1 H, H6a), 3.88(d,  $J = 7.0$  Hz, 1 H, H3), 4.16(d,  $J = 8.3$  Hz, 1 H, H20b), 4.27(br s, 2 H,  $\text{ClCH}_2$ ), 4.31(d,  $J = 8.3$  Hz, 1 H, H20a), 4.44(ddd,  $J = 10.9, 6.7, 4.5$  Hz, 1 H, H7), 4.90(m, 1 H, H13), 4.98(dd,  $J = 9.7, 2.1$  Hz, 1 H, H5), 5.64(d,  $J = 7.0$  Hz, 1 H, H2), 6.41(s, 1 H, H10), 7.49(dd,  $J = 7.9, 7.4$  Hz, 2 H, benzoate, *m*), 7.61(tt,  $J = 7.4, 1.3$  Hz, 1 H, benzoate, *p*), 8.11(d,  $J = 7.9, 1.3$  Hz, 2 H, benzoate, *o*) ppm.  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ )  $\delta$  9.3(Me(19)), 15.3(Me(18)), 20.6(4-Ac), 22.3, 26.7(Me16, Me17), 35.8(C(6)), 38.6(C(14)), 40.5( $\text{ClCH}_2$ ), 42.6(C(15)), 46.2(C(3)), 58.8(C(8)), 68.0(C(13)), 72.0, 75.0, 75.9(C(7), C(2), C(10), C(20)), 79.0(C(1)), 80.9(C(4)), 84.4(C(5)), 128.8, 129.7, 130.3, 133.9(benzoate), 131.8(C(11)), 147.1(C(12)), 167.4, 167.7 ( $\text{ClCH}_2\text{C(O)O}$ , benzoate), 171.0 (4-Ac), 203.7(C(9))ppm. Anal. Calcd for  $\text{C}_{31}\text{H}_{37}\text{ClO}_{11}$ .  $\text{H}_2\text{O}$ : C, 58.26; H, 6.15. Found: C, 58.26; H, 6.07.

**[0062] 10-Propionyl-10-DAB.** To a solution of 10-DAB (47 mg, 0.086 mmol) in THF (2 mL) at room temperature was added a mixture of propionic anhydride (4 mL) and  $\text{ZnCl}_2/\text{THF}$  solution (0.5 M, 350 mL, 0.173 mmol, 2 equiv) under  $\text{N}_2$ . The reaction mixture was stirred at room temperature for 14 h. Then the reaction mixture was diluted with EtOAc (150 mL), exhaustively washed with saturated aqueous  $\text{NaHCO}_3$  solution (50 mLx3), brine. The organic layer was dried over  $\text{Na}_2\text{SO}_4$ , concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:1) as the eluent and dried in *vacuo* to give 10-propionyl-10-DAB as a white solid: yield, 48 mg (93%). mp 212-213 °C dec;  $[\alpha]_{\text{D}}^{25} -96^\circ$  ( $\text{CHCl}_3$ ,  $c = 0.78$ );  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )  $\delta$  1.11(s, 6 H, Me16, Me17), 1.24 (t,  $J = 7.6$  Hz, 3 H,  $\text{CH}_3\text{CH}_2$ ), 1.60(s, 1 H, 1-OH), 1.67(s, 3 H, Me19), 1.87(ddd,  $J = 14.7, 10.9, 2.2$  Hz, 1 H, H6b), 2.05 (d,  $J = 5.1$  Hz, 1 H, 13-OH), 2.06(d,  $J = 1.3$  Hz, 3 H, Me18), 2.28(s, 3 H, 4-Ac), 2.30(d,  $J = 7.5$  Hz, 2 H, H14a, H14b), 2.51(d,  $J = 4.1$  Hz, 1 H, 7-OH), 2.55(q,  $J = 7.6$  Hz, 2 H,  $\text{CH}_3\text{CH}_2$ ), 2.57(ddd,  $J = 14.7, 9.5, 6.7$  Hz, 1 H, H6a), 3.90(d,  $J = 6.9$  Hz, 1 H, H3), 4.16(dd,  $J = 8.4, 0.8$  Hz, 1 H, H20b), 4.31(d,  $J = 8.4$  Hz, 1 H, H20a), 4.48(ddd,  $J = 10.9, 6.7, 4.1$  Hz, 1 H, H7), 4.90(m, 1 H, H13), 4.99(dd,  $J = 9.5, 2.2$  Hz, 1 H, H5), 5.63(d,  $J = 6.9$  Hz, 1 H, H2), 6.34(s, 1 H, H10), 7.48(dd,  $J = 8.1, 7.4$  Hz, 2 H, benzoate, *m*), 7.61(tt,  $J = 7.4, 1.3$  Hz, 1 H, benzoate, *p*), 8.11(dd,  $J = 8.3, 1.3$  Hz, 2 H, benzoate, *o*) ppm.  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ )  $\delta$  8.8 ( $\text{CH}_3\text{CH}_2$ ), 9.2(Me(19)), 15.2(Me(18)), 20.7(4-Ac), 22.3, 26.8, 27.4(Me16, Me17,  $\text{CH}_3\text{CH}_2$ ), 35.5(C(6)), 38.7(C(14)), 42.6(C(15)), 46.1(C(3)), 58.7(C(8)), 67.9(C(13)), 72.3, 75.1, 76.1, 76.5(C(7), C(2), C(10), C(20)), 79.1(C(1)), 80.9(C(4)), 84.5(C(5)), 128.7, 129.7, 130.3, 133.8(benzoate), 132.3(C(11)), 146.5(C(12)), 167.4(benzoate), 170.9, 174.9(4-Ac, 10-C(O)O), 204.6(C(9))ppm. Anal. Calcd for  $\text{C}_{32}\text{H}_{40}\text{O}_{11}$ : C, 63.99; H, 6.71. Found: C, 63.81; H, 6.80.

#### C. Selective acylation of a C(10) hydroxyl group using $\text{CeCl}_3$ :

**[0063] General procedure:** To a solution of 10-DAB in THF (20 mL per mmol of 10-DAB) under  $\text{N}_2$  was added  $\text{CeCl}_3$  and the appropriate anhydride or pyrocarbonate (amounts specified in Table 1). The reaction mixture was stirred at 25 °C and monitored by TLC analysis. When that analysis indicated complete reaction (time specified in Table 1), the reaction mixture was diluted with EtOAc and washed three times with saturated aqueous sodium bicarbonate solution. The combined bicarbonate washings were extracted three times with EtOAc, the organic layers were combined and dried over sodium sulfate, and the solvent was evaporated. The crude product was purified by flash column chromatography. Further purification, if necessary, was carried out by recrystallization from EtOAc/Hexane.

Table 1. CeCl<sub>3</sub> catalyzed acylation of 10-DAB

$  \begin{array}{c}  \text{10-DAB} \xrightarrow[\text{THF, RT}]{(\text{RCO})_2\text{O, CeCl}_3} \text{10-Acyl-10-DAB}  \end{array}  $				
Entry	R (eq)	CeCl <sub>3</sub> (eq)	Time (hr)	Yield (%)
1	Me (10)	0.1	1.5	91
2	Pr (10)	0.1	3	100
3	iPr (10)	0.1	4.5	100
4	Ph (10)	0.1	21	94
5	cyclopropyl (10)	0.1	20.5	94
6	MeCH=CH (10)	0.1	20	91
7	CH <sub>2</sub> =CHCH <sub>2</sub> O (5)	0.1	1	96
8	EtO (5)	0.1	3	99
9	MeO (5)	0.1	3	98
10	tBuO (10)	0.7	24	94
11	BnO (3)	0.7	1	98

[0064] 10-butyl-10-DAB. mp 145-149 °C;  $[\alpha]_{\text{D}}^{25}$  -86.6 (CHCl<sub>3</sub>, c = 1); <sup>1</sup>H NMR (500 MHz, CDCl<sub>3</sub>) δ 8.13-8.11 (2H, m), 7.62 (1H, m), 7.51-7.48 (2H, m), 6.35 (1H, s), 5.64 (1H, d, J 7.0Hz), 4.99 (1H, d, J 7.7Hz), 4.90 (1H, m), 4.48 (1H, m), 4.31 (1H, d, J 8.3Hz), 4.18 (1H, d, J 8.3Hz), 3.91 (1H, d, J 7.0Hz), 2.60-2.42 (4H, m), 2.36-2.26 (2H, m), 2.28 (3H, s), 2.06 (3H, d, J 1.0Hz), 1.88 (1H, ddd, J 1.9, 10.9, 13.0Hz), 1.76 (2H, hex, J 7.4Hz), 1.68 (3H, s), 1.12 (6H, s) and 1.04 (3H, t, J 7.4Hz); <sup>13</sup>C NMR (100MHz, CDCl<sub>3</sub>) δ 204.2, 173.9, 170.6, 167.1, 146.2, 133.7, 132.0, 130.1, 129.4, 128.6, 84.5, 88.9, 79.1, 76.5, 76.0, 75.0, 72.3, 68.0, 58.8, 46.2, 42.7, 38.7, 37.1, 36.2, 35.6, 30.6, 27.0, 22.6, 20.9, 18.4, 17.8, 15.5 and 9.4; Anal. Calcd. for C<sub>33</sub>H<sub>42</sub>O<sub>11</sub>: C, 64.48; H, 6.89 Found: C, 63.67; H, 7.01.

[0065] 10-isobutyl-10-DAB. mp 143 °C;  $[\alpha]_{\text{D}}^{25}$  -62.6° (CHCl<sub>3</sub>, c=0.075); <sup>1</sup>H NMR (CDCl<sub>3</sub>, 500MHz): δ 8.12 (2H, d, J 7.3Hz), 7.62 (1H, m), 7.51-7.48 (2H, m), 6.33 (1H, s), 5.65 (1H, d, J 7.3Hz), 5.00 (1H, d, J 7.9Hz), 4.91 (1H, m), 4.48 (1H, ddd, J 4.3, 6.7, 11.0Hz), 4.31 (1H, d, J 8.6Hz), 4.18 (1H, d, J 8.6Hz), 3.91 (1H, d, J 7.3Hz), 2.74 (1H, pent, J 6.7Hz), 2.57 (1H, m), 2.51 (1H, d, J 4.3Hz), 2.31 (1H, m), 2.28 (3H, s), 2.06 (3H, s), 2.01 (1H, d, J 5.5Hz), 1.90 (1H, ddd, J 2.3, 11.0, 14.6Hz), 1.68 (3H, s), 1.60 (1H, s), 1.51 (3H, s), 1.33 (3H, d, J 6.7Hz), 1.26 (3H, d, J 6.7Hz), 1.13 (3H, s) and 1.12 (3H, s); <sup>13</sup>C NMR (100MHz, CDCl<sub>3</sub>) δ 204.1, 177.2, 170.6, 167.1, 146.2, 133.7, 132.1, 130.1, 129.4, 128.6, 95.5, 84.5, 80.9, 79.1, 76.5, 75.8, 74.9, 72.3, 68.0, 58.8, 46.2, 42.7, 38.7, 35.6, 34.1, 27.0, 22.6, 20.9, 19.2, 18.7, 15.5 and 9.4; Anal. Calcd. for C<sub>33</sub>H<sub>42</sub>O<sub>11</sub>·0.5H<sub>2</sub>O: C, 64.48; H, 6.89 Found: C, 63.05; H, 6.70.

[0066] 10-benzoyl-10-DAB. <sup>1</sup>H NMR (CDCl<sub>3</sub>, 500MHz): δ 8.15-8.11 (4H, m), 7.64-7.6 (2H, m), 7.52-7.48 (4H, m), 6.62 (1H, s), 5.7 (1H, d, J 7.1Hz), 5.02 (1H, d, J 7.7Hz), 4.94 (1H, m), 4.57 (1H, ddd, J 4.4, 7.1, 11.0Hz), 4.33 (1H, d, J 8.2Hz), 4.20 (1H, d, J 8.3Hz), 3.99 (1H, d, J 6.6Hz), 2.62 (1H, ddd, J 6.6, 9.3, 14.8), 2.55 (1H, d, J 4.4Hz), 2.35 (2H, m), 2.30 (3H, s), 2.13 (3H, d, J 1.1Hz), 2.03 (1H, d, J 4.9Hz), 1.91 (1H, ddd, J 2.2, 11.0, 13.2Hz), 1.71 (3H, s), 1.65 (1H, s), 1.25 (3H, s) and 1.21 (3H, s); <sup>13</sup>C NMR (100MHz, CDCl<sub>3</sub>) δ 204.0, 170.7, 167.1, 166.5, 146.5, 133.7, 133.6, 132.0, 130.1, 129.9, 129.4, 129.3, 128.7, 128.5, 84.5, 80.9, 79.1, 76.5, 75.0, 72.4, 68.1, 58.8, 46.3, 42.8, 38.7, 35.8, 29.7, 27.2, 22.6, 21.2, 15.6 and 9.5; Anal. Calcd. for C<sub>35</sub>H<sub>40</sub>O<sub>11</sub>: C, 66.66; H, 6.22 Found C, 66.46; H, 6.19.

[0067] 10-trans crotonyl-10-DAB. <sup>1</sup>H NMR (CDCl<sub>3</sub>, 500MHz): δ 8.13 (2H, d, J 7.1Hz), 7.62 (1H, m), 7.51-7.48 (2H, m), 7.11 (1H, m), 6.42 (1H, s), 6.02 (1H, dq, J 1.7, 15.4Hz), 5.66 (1H, d, J 7.1Hz), 4.99 (1H, dd, J 2.0, 9.6Hz), 4.91 (1H, t, J 7.6Hz), 4.50 (1H, dd, J 7.1, 10.8Hz), 4.31 (1H, d, J 8.3Hz), 4.19 (1H, d, J 8.3Hz), 3.93 (1H, d, J 7.1Hz), 2.61-2.55 (2H, m), 2.33-2.31 (2H, m), 2.28 (3H, s), 2.07 (3H, d, J 1.5Hz), 1.95 (3H, dd, J 1.6, 6.8Hz), 1.89 (1H, ddd, J 2.3, 11.0, 13.4Hz), 1.68 (3H, s), 1.15 (3H, s) and 1.14 (3H, s); <sup>13</sup>C NMR (75MHz, CDCl<sub>3</sub>) δ 212.4, 181.0, 170.8, 167.3, 166.5, 146.4, 133.8, 132.3, 130.2, 129.5, 128.7, 121.9, 116.0, 84.7, 84.6, 80.9, 79.2, 77.2, 75.9, 75.1, 72.4, 68.1, 58.8, 46.1, 42.7, 38.6, 35.6, 27.0, 20.9, 18.0, 15.4 and 9.3; Anal. Calcd. for C<sub>33</sub>H<sub>40</sub>O<sub>11</sub>: C, 64.69; H, 6.58 Found C, 63.93; H, 6.61.

[0068] 10-cyclopropanoyl-10-DAB.  $^1\text{H}$  ( $\text{CDCl}_3$ , 500MHz):  $\delta$  8.12 (2H, d,  $J$  7.3Hz), 7.62 (1H, t,  $J$  7.5Hz), 7.49 (2H, t,  $J$  7.7Hz), 6.35 (1H, s), 5.65 (1H, d,  $J$  7.0Hz), 4.99 (1H, app-d,  $J$  8.2Hz), 4.91 (1H, m), 4.46 (1H, ddd,  $J$  4.1, 6.8, 10.8Hz), 4.31 (1H, d,  $J$  8.1Hz), 4.18 (1H, d,  $J$  8.1Hz), 3.90 (1H, d,  $J$  7.0Hz), 2.56 (1H, m), 2.51 (1H, d,  $J$  4.1Hz), 2.31 (2H, m), 2.07 (3H, d,  $J$  1.0Hz), 2.00 (1H, d,  $J$  4.9Hz), 1.87 (1H, ddd,  $J$  2.1, 10.8, 14.6Hz), 1.79 (1H, ddd,  $J$  3.4, 7.9, 12.4Hz), 1.68 (3H, s), 1.60 (1H, s), 1.16-1.14 (2H, m), 1.13 (6H, s) and 1.01-0.97 (2H, m);  $^{13}\text{C}$  NMR (100MHz,  $\text{CDCl}_3$ )  $\delta$  204.3, 175.2, 170.6, 167.1, 146.4, 133.7, 132.0, 130.1, 129.4, 128.6, 84.5, 80.9, 79.1, 76.5, 76.0, 74.9, 72.4, 68.0, 58.8, 46.2, 42.7, 38.6, 35.6, 34.0, 27.0, 25.6, 24.9, 22.6, 21.0, 15.6, 13.1, 9.4 and 9.1; Anal. Calcd. for  $\text{C}_{33}\text{H}_{40}\text{O}_{11}$ : C, 64.69; H, 6.58 Found: C, 64.47; H, 6.66.

[0069] 10-Ethoxycarbonyl-10-DAB. mp 214-215 °C;  $[\alpha]_{\text{D}}^{25}$  -81° ( $\text{CHCl}_3$ ,  $c$  = 0.35);  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  1.13 (s, 3 H, Me17), 1.14 (s, 3 H, Me16), 1.38 (t,  $J$  = 7.1 Hz, 3 H,  $\text{CH}_3\text{CH}_2$ ), 1.59 (s, 1 H, 1-OH), 1.70 (s, 3 H, Me19), 1.88 (ddd,  $J$  = 14.6, 10.5, 2.1 Hz, 1 H, H6b), 2.00 (d,  $J$  = 5.0 Hz, 1 H, 13-OH), 2.10 (d,  $J$  = 1.4 Hz, 3 H, Me18), 2.28 (s, 3 H, 4-Ac), 2.30 (m, 2 H, H14a, H14b), 2.46 (d,  $J$  = 4.2 Hz, 1 H, 7-OH), 2.57 (ddd,  $J$  = 14.6, 9.6, 6.7 Hz, 1 H, H6a), 3.88 (d,  $J$  = 6.9 Hz, 1 H, H3), 4.18 (d,  $J$  = 8.2 Hz, 1 H, H20b), 4.31 (d,  $J$  = 8.2 Hz, 1 H, H20a), 4.23-4.33 (m, 2 H,  $\text{CH}_3\text{CH}_2$ ), 4.44 (ddd,  $J$  = 10.5, 6.7, 4.2 Hz, 1 H, H7), 4.90 (m, 1 H, H13), 4.98 (dd,  $J$  = 9.6, 2.1 Hz, 1 H, H5), 5.65 (d,  $J$  = 6.9 Hz, 1 H, H2), 6.17 (s, 1 H, H10), 7.48 (dd,  $J$  = 8.2, 7.3 Hz, 2 H, benzoate, *m*), 7.60 (tt,  $J$  = 7.3, 1.4 Hz, 1 H, benzoate, *p*), 8.11 (d,  $J$  = 8.2, 1.4 Hz, 2 H, benzoate, *o*) ppm;  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ )  $\delta$  9.2, 14.0, 15.5, 20.8, 22.4, 26.7, 35.4, 38.5, 42.4, 46.0, 58.6, 65.0, 67.7, 72.2, 74.9, 76.4, 78.7, 79.0, 80.6, 84.4, 128.7, 129.4, 130.1, 131.5, 133.7, 147.5, 155.4, 167.1, 170.8, 204.7 ppm.

[0070] 10-Methoxycarbonyl-10-DAB. mp 218-219 °C;  $[\alpha]_{\text{D}}^{25}$  -83° ( $\text{CHCl}_3$ ,  $c$  = 0.58);  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  1.12 (s, 3 H, Me17), 1.13 (s, 3 H, Me16), 1.59 (s, 1 H, 1-OH), 1.70 (s, 3 H, Me19), 1.88 (ddd,  $J$  = 14.7, 10.8, 1.8 Hz, 1 H, H6b), 2.00 (d,  $J$  = 5.0 Hz, 1 H, 13-OH), 2.10 (d,  $J$  = 1.4 Hz, 3 H, Me18), 2.28 (s, 3 H, 4-Ac), 2.30 (m, 2 H, H14a, H14b), 2.40 (d,  $J$  = 4.1 Hz, 1 H, 7-OH), 2.57 (ddd,  $J$  = 14.7, 9.7, 6.6 Hz, 1 H, H6a), 3.87 (d,  $J$  = 6.9 Hz, 1 H, H3), 3.88 (s, 3 H, MeOC(O)), 4.18 (d,  $J$  = 8.4 Hz, 1 H, H20b), 4.31 (d,  $J$  = 8.4 Hz, 1 H, H20a), 4.44 (ddd,  $J$  = 10.8, 6.6, 4.1 Hz, 1 H, H7), 4.90 (m, 1 H, H13), 4.98 (dd,  $J$  = 9.7, 1.8 Hz, 1 H, H5), 5.65 (d,  $J$  = 6.9 Hz, 1 H, H2), 6.17 (s, 1 H, H10), 7.48 (t,  $J$  = 8.2, 7.3 Hz, 2 H, benzoate, *m*), 7.61 (tt,  $J$  = 7.3, 1.4 Hz, 1 H, benzoate, *p*), 8.11 (d,  $J$  = 8.2, 1.4 Hz, 2 H, benzoate, *o*) ppm;  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ )  $\delta$  9.2, 15.5, 20.7, 22.4, 26.7, 35.5, 38.5, 42.4, 46.0, 55.4, 58.6, 65.0, 67.7, 72.1, 74.8, 76.4, 78.9, 79.0, 80.6, 84.4, 128.7, 129.4, 130.1, 131.4, 133.7, 147.5, 155.9, 167.1, 170.8, 204.6 ppm.

[0071] 10-*t*Boc-10-DAB. mp 193-194 °C;  $[\alpha]_{\text{D}}^{25}$  -82° ( $\text{CHCl}_3$ ,  $c$  = 0.33);  $^1\text{H}$  NMR (500 MHz,  $\text{CDCl}_3$ )  $\delta$  1.13 (s, 6 H, Me17, Me16), 1.48 (s, 9 H, *t*BuO), 1.58 (s, 1 H, 1-OH), 1.69 (s, 3 H, Me19), 1.88 (ddd,  $J$  = 14.9, 11.0, 2.2 Hz, 1 H, H6b), 1.99 (d,  $J$  = 5.0 Hz, 1 H, 13-OH), 2.08 (d,  $J$  = 1.4 Hz, 3 H, Me18), 2.28 (s, 3 H, 4-Ac), 2.30 (m, 2 H, H14a, H14b), 2.56 (ddd,  $J$  = 14.9, 9.6, 6.9 Hz, 1 H, H6a), 2.68 (d,  $J$  = 3.6 Hz, 1 H, 7-OH), 3.88 (d,  $J$  = 6.9 Hz, 1 H, H3), 4.19 (d,  $J$  = 8.2 Hz, 1 H, H20b), 4.31 (d,  $J$  = 8.2 Hz, 1 H, H20a), 4.46 (ddd,  $J$  = 11.0, 6.9, 3.6 Hz, 1 H, H7), 4.90 (m, 1 H, H13), 4.99 (dd,  $J$  = 9.6, 2.2 Hz, 1 H, H5), 5.64 (d,  $J$  = 6.9 Hz, 1 H, H2), 6.11 (s, 1 H, H10), 7.48 (t,  $J$  = 7.8 Hz, 2 H, benzoate, *m*), 7.60 (tt,  $J$  = 7.8, 1.3 Hz, 1 H, benzoate, *p*), 8.11 (dd,  $J$  = 7.8, 1.3 Hz, 2 H, benzoate, *o*) ppm;  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ )  $\delta$  9.2, 15.6, 20.9, 22.4, 26.8, 27.5, 35.3, 38.5, 42.5, 45.9, 58.7, 67.9, 72.3, 74.7, 76.4, 78.0, 79.2, 80.8, 83.8, 84.5, 128.7, 129.4, 130.1, 131.8, 133.7, 147.3, 154.0, 167.2, 170.8, 205.0 ppm.

#### D. Selective carbamoylation of a C(10) hydroxyl group:

[0072] General procedure for the Selective Carbamoylation of the C-10 Hydroxyl group of 10-DAB: A solution of 0.061 mmol (1.1 mol equiv) of the isocyanate in 2 mL of THF was added, under nitrogen, to a mixture of 10-DAB (30 mg, 0.055 mmol) and CuCl (5.5 mg, 0.055 mmol) at 0 °C. The mixture was stirred for the time indicated in Table 2. After this time the reaction was warmed to 25 °C and stirring was continued for the time indicated in Table 2. The reaction was quenched by the addition of saturated aqueous ammonium chloride solution, and the mixture was extracted three times with EtOAc. The combined organic layers were washed with saturated aqueous  $\text{NaHCO}_3$  solution, dried over sodium sulfate, and the solvent was evaporated to yield a white solid. The product was purified by flash column chromatography using 2:1 EtOAc/hexane as the eluent.

Table 2. Carbamoylation of 10-DAB

10-DAB $\xrightarrow[\text{THF}]{\text{RNCO, CuCl}}$ 10-Carbamoyl-10-DAB				
Entry	R (eq)	Temp (°C)	Time (hr)	Yield (%)
1	Et (1.1)	0 rt	7.5 0.5	88
2	allyl (1.1)	0 rt	6 0.5	88
3	Bu (1.1)	0 rt	6.5 0.5	87
4	Ph (1.1)	rt	3	94

[0073] 10-ethylcarbamoyl-10-DAB. mp 241-243 °C;  $[\alpha]_{\text{H}_g} -92.0^\circ$  ( $\text{CHCl}_3$ ,  $c=0.5$ );  $^1\text{H}$  NMR (400MHz,  $\text{CDCl}_3$ )  $\delta$  8.13 (2H, d,  $J$  7.1Hz), 7.63 (1H, m), 7.52-7.48 (2H, m), 6.27 (1H, s), 5.63 (1H, d,  $J$  6.9Hz), 5.01 (1H, dd,  $J$  1.9, 9.6Hz), 4.97 (1H, m), 4.91 (1H, m), 4.50 (1H, ddd,  $J$  3.7, 6.5, 10.5Hz), 4.31 (1H, d,  $J$  8.3Hz), 4.17 (1H, d,  $J$  8.3Hz), 3.88 (1H, d,  $J$  7.0Hz), 3.32-3.25 (2H, m), 3.10 (1H, d,  $J$  3.7Hz), 2.56 (1H, ddd,  $J$  6.8, 9.8, 14.8Hz), 2.31 (1H, m), 2.29 (3H, s), 2.09 (3H, s), 1.88 (1H, ddd,  $J$  2.2, 11.0, 13.3Hz), 1.67 (3H, s), 1.60 (1H, s), 1.19 (3H, t,  $J$  7.2Hz) and 1.10 (6H, s); Anal. Calcd. for  $\text{C}_{32}\text{H}_{40}\text{NO}_{11}$ : C, 62.43; H, 6.71 Found: C, 61.90; H, 6.77.

[0074] 10-butylcarbamoyl-10-DAB.  $[\alpha]_{\text{H}_g} -89.6^\circ$  ( $\text{CHCl}_3$ ,  $c=0.25$ );  $^1\text{H}$  NMR (500MHz,  $\text{CDCl}_3$ )  $\delta$  8.12 (2H, d,  $J$  7.3Hz), 7.61 (1H, m), 7.51-7.45 (2H, m), 6.27 (1H, s), 5.64 (1H, d,  $J$  6.7Hz), 5.00 (1H, d,  $J$  8.0Hz), 4.91 (1H, m), 4.49 (1H, m), 4.31 (1H, d,  $J$  8.5Hz), 4.19 (1H, d,  $J$  8.5Hz), 3.89 (1H, d,  $J$  6.7Hz), 3.25-3.23 (2H, m), 3.04 (1H, m), 2.56 (1H, ddd,  $J$  6.7, 9.7, 14.7Hz), 2.30 (1H, d,  $J$  7.9Hz), 2.28 (3H, s), 2.09 (3H, s), 1.99 (1H, d,  $J$  4.9Hz), 1.88 (1H, ddd,  $J$  2.5, 11.0, 13.4Hz), 1.68 (3H, s), 1.59 (1H, s), 1.55 (2H, b), 1.42-1.37 (2H, m), 1.11 (6H, s) and 0.95 (3H, t,  $J$  7.6Hz); Anal. Calcd. for  $\text{C}_{34}\text{H}_{44}\text{NO}_{11}$ : C, 63.44; H, 7.05 Found: C, 62.64; H, 7.01.

[0075] 10-phenylcarbamoyl-10-DAB. mp 178-180 °C;  $[\alpha]_{\text{H}_g} -93.0^\circ$  ( $\text{CHCl}_3$ ,  $c=0.5$ );  $^1\text{H}$  NMR (400Hz,  $\text{CDCl}_3$ )  $\delta$  8.13 (2H, d,  $J$  6.9Hz), 7.63 (1H, t,  $J$  7.4Hz), 7.51 (2H, t,  $J$  7.6Hz), 7.42 (1H, d,  $J$  7.8Hz), 7.36-7.32 (2H, m), 7.12 (1H, t,  $J$  7.4Hz), 6.87 (1H, b), 6.38 (1H, s), 5.66 (1H, d,  $J$  7.0Hz), 5.02 (1H, app d,  $J$  7.8Hz), 5.93 (1H, m), 4.52 (1H, ddd,  $J$  3.8, 6.5, 10.5Hz), 4.33 (1H, d,  $J$  8.3Hz), 4.18 (1H, d,  $J$  8.3Hz), 3.91 (1H, d,  $J$  7.0Hz), 2.83 (1H, d,  $J$  4.0Hz), 2.59 (1H, ddd,  $J$  6.5, 9.4, 14.5Hz), 2.33 (1H, m), 2.29 (3H, s), 2.12 (3H, d,  $J$  1.4Hz), 2.04 (1H, d,  $J$  5.1Hz), 1.89 (1H, ddd,  $J$  2.2, 11.0, 14.4Hz), 1.69 (3H, s), 1.62 (1H, s), 1.15 (3H, s) and 1.13 (3H, s).

[0076] 10-allylcarbamoyl-10-DAB. mp 165-170 °C;  $[\alpha]_{\text{H}_g} -80.0^\circ$  ( $\text{CHCl}_3$ ,  $c=0.25$ );  $^1\text{H}$  NMR (500MHz,  $\text{CDCl}_3$ )  $\delta$  8.12 (2H, d,  $J$  7.3Hz), 7.62 (1H, m), 7.51-7.48 (2H, m), 6.27 (1H, s), 5.89 (1H, m), 5.62 (1H, d,  $J$  6.7Hz), 5.31 (1H, s), 5.19 (1H, d,  $J$  9.8Hz), 5.08 (1H, m), 5.00 (1H, d,  $J$  7.9Hz), 4.90 (1H, m), 4.49 (1H, ddd,  $J$  3.7, 6.1, 10.4Hz), 4.31 (1H, d,  $J$  8.5Hz), 4.17 (1H, d,  $J$  8.5Hz), 3.88-3.86 (2H, m), 3.03 (1H, d,  $J$  3.7Hz), 2.55 (1H, ddd,  $J$  6.7, 9.8, 15.9Hz), 2.30 (1H, m), 2.29 (3H, s), 2.08 (3H, s), 2.06 (1H, app d,  $J$  4.9Hz), 1.87 (1H, ddd,  $J$  1.8, 11.0, 14.0Hz), 1.67 (3H, s), 1.58 (1H, s) and 1.09 (6H, s); Anal. Calcd. for  $\text{C}_{33}\text{H}_{40}\text{NO}_{11}$ : C, 63.15; H, 6.58 Found: C, 61.73; H, 6.45.

#### Example 2

[0077] General Procedure for the Preparation of 7-silyl-10-TES-10-DAB. To a solution of 7-triethylsilyl-10-DAB, 7-*t*-butyldimethylsilyl-10-DAB, or 7-dimethylisopropylsilyl-10-DAB in THF at 0 °C was slowly added N, O-bis(triethylsilyl) trifluoroacetamide (5 equiv), and a catalytic amount of LiHMDS/THF solution (5 mol %), respectively, under  $\text{N}_2$ . The reaction mixture was stirred at 0 °C for 15 min. EtOAc (10 mL) was added, and the solution was filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:2) as the eluent and dried in *vacuo* overnight to give, respectively, 7,10-bis(triethylsilyl)-10-DAB (95% yield), 7-*t*-butyldimethylsilyl-10-triethylsilyl-10-DAB (98% yield), or 7-dimethylisopropylsilyl-10-triethylsilyl-10-DAB (94% yield).

[0078] 7-Dimethylphenylsilyl-10-TBS-10-DAB. To a solution of 7-dimethylphenylsilyl-10-DAB (35 mg, 0.052 mmol) in THF (2 mL) at 0 °C was added N, O-bis(*t*-butyldimethylsilyl)trifluoroacetamide (337  $\mu\text{L}$ , 1.09 mmol, 20 equiv), and a catalytic amount of LiHMDS/THF solution (1 M, 6  $\mu\text{L}$ , 0.006 mmol), respectively, under  $\text{N}_2$ . The reaction mixture was

stirred at 0 °C for 4 h, then warmed to room temperature for an additional 4 h. EtOAc (10 mL) was added, and the solution was filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:2) as the eluent and dried *in vacuo* overnight to give 39 mg (92% yield) of 7-dimethylphenylsilyl-10-t-butyldimethylsilyl-10-DAB.

### Example 3

#### Selective silylation of a C(7) hydroxyl group

**[0079] 7-TBS-10-DAD.** To a mixture of 10-DAB (38 mg, 0.070 mmol), imidazole (190 mg, 2.79 mmol, 40 equiv), and tert-butyldimethylsilyl chloride (210 mg, 1.40 mmol, 20 equiv) was added DMF (0.1 mL) at room temperature under N<sub>2</sub>. The reaction mixture was vigorously stirred at room temperature for 24 h. EtOAc (20 mL) was added, and the solution was filtered through a short column of silica gel. The silica gel was washed with EtOAc (200 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using 10% EtOAc-CH<sub>2</sub>Cl<sub>2</sub> as the eluent and dried *in vacuo* overnight to give 7-TBS-10-DAB as a white solid: yield, 41 mg (90%). mp 222-223 °C; [α]<sub>D</sub><sup>20</sup> -51° (CHCl<sub>3</sub>, c = 0.36); <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 0.05, 0.06 (2 s, 6 H, Me<sub>2</sub>Si), 0.83 (s, 9 H, Me<sub>3</sub>C), 1.09 (s, 6 H, Me16, Me17), 1.57 (s, 1 H, 1-OH), 1.75 (s, 3 H, Me19), 1.87 (ddd, J = 14.4, 10.6, 2.0 Hz, 1 H, H6b), 2.01 (d, J = 5.0 Hz, 1 H, 13-OH), 2.09 (d, J = 1.3, 3 H, Me18), 2.28 (m, 2 H, H14a, H14b), 2.29 (s, 3 H, 4-Ac), 2.46 (ddd, J = 14.4, 9.6, 6.7 Hz, 1 H, H6a), 3.96 (d, J = 6.9 Hz, 1 H, H3), 4.16 (d, J = 8.3 Hz, 1 H, H20b), 4.24 (d, J = 2.2 Hz, 1 H, 10-OH), 4.31 (d, J = 8.3 Hz, 1 H, H20a), 4.38 (dd, J = 10.6, 6.7 Hz, 1 H, H7), 4.88 (m, 1 H, H13), 4.96 (dd, J = 9.6, 2.0 Hz, 1 H, H5), 5.15 (d, J = 2.0 Hz, 1 H, H10), 5.60 (d, J = 6.9 Hz, 1 H, H2), 7.47 (dd, J = 8.1, 7.5 Hz, 2 H, benzoate, m), 7.60 (tt, J = 7.5, 1.3 Hz, 1 H, benzoate, p), 8.10 (d, J = 8.1, 1.3 Hz, 2 H, benzoate, o) ppm. <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>) δ -5.8, -3.8 (Me<sub>2</sub>Si), 9.7 (Me(19)), 14.8 (Me(18)), 17.6 (Me<sub>3</sub>C), 19.3 (4-Ac), 22.4, 26.7 (Me16, Me17), 25.4 (Me<sub>3</sub>C), 37.4 (C(6)), 38.7 (C(14)), 42.7 (C(15)), 47.0 (C(3)), 58.0 (C(8)), 68.0 (C(13)), 73.1, 74.7, 75.0 (C(7), C(2), C(10), C(20)), 78.9 (C(1)), 80.9 (C(4)), 84.3 (C(5)), 128.8, 129.8, 130.3, 133.8 (benzoate), 135.7 (C(11)), 141.9 (C(12)), 167.4 (benzoate), 171.2 (4-Ac), 210.8 (C(9)) ppm. Anal. Calcd for C<sub>35</sub>H<sub>50</sub>O<sub>10</sub>Si: C, 63.80; H, 7.65. Found: C, 63.72; H, 7.70.

**[0080] 7-Dimethylphenylsilyl-10-DAB.** To a THF (3 mL) solution of 10-DAB (54 mg, 0.099 mmol) at -20 °C was added pyridine (0.6 mL), dimethylphenylsilyl chloride (250 mL, 1.49 mmol, 15 equiv) under N<sub>2</sub>. The reaction mixture was stirred at -20 °C for 2 h. EtOAc (10 mL) and saturated NaHCO<sub>3</sub> aqueous solution (0.5 mL) was added, and the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: CH<sub>2</sub>Cl<sub>2</sub> (1:10) as the eluent and dried overnight *in vacuo* to give 7-dimethylphenylsilyl-10-DAB as a white solid: yield, 62 mg (92%). mp 219-220 °C; [α]<sub>D</sub><sup>20</sup> -28° (CHCl<sub>3</sub>, c = 0.27); <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 0.35, 0.37 (2 s, 6 H, Me<sub>2</sub>Si), 1.05 (s, 3 H, Me17), 1.06 (s, 3 H, Me16), 1.54 (s, 1 H, 1-OH), 1.73 (d, J = 1.1, 3 H, Me18), 1.76 (s, 3 H, Me19), 1.90 (ddd, J = 14.4, 10.6, 2.1 Hz, 1 H, H6b), 1.93 (d, J = 5.0 Hz, 1 H, 13-OH), 2.23 (m, 2 H, H14a, H14b), 2.25 (s, 3 H, 4-Ac), 2.43 (ddd, J = 14.4, 9.6, 6.8 Hz, 1 H, H6a), 3.86 (d, J = 7.0 Hz, 1 H, H3), 4.10 (d, J = 2.1 Hz, 1 H, 10-OH), 4.16 (d, J = 8.3 Hz, 1 H, H20b), 4.28 (d, J = 8.3 Hz, 1 H, H20a), 4.31 (dd, J = 10.6, 6.8 Hz, 1 H, H7), 4.81 (m, 1 H, H13), 4.84 (d, J = 2.1 Hz, 1 H, H10), 4.90 (dd, J = 9.6, 2.1 Hz, 1 H, H5), 5.59 (d, J = 7.0 Hz, 1 H, H2), 7.41, 7.53 (2 m, 5 H, C<sub>6</sub>H<sub>5</sub>), 7.46 (dd, J = 8.0, 7.5 Hz, 2 H, benzoate, m), 7.55 (tt, J = 7.5, 1.2 Hz, 1 H, benzoate, p), 8.09 (d, J = 8.0, 1.2 Hz, 2 H, benzoate, o) ppm. <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>) δ -1.8, -1.1 (Me<sub>2</sub>Si), 9.8 (Me(19)), 14.4 (Me(18)), 19.4 (4-Ac), 22.3, 26.7 (Me16, Me17), 37.2 (C(6)), 38.6 (C(14)), 42.6 (C(15)), 46.7 (C(3)), 58.0 (C(8)), 68.0 (C(13)), 73.2, 74.7, 75.0 (C(7), C(2), C(10), C(20)), 78.8 (C(1)), 80.8 (C(4)), 84.3 (C(5)), 128.3, 128.8, 129.8, 130.2, 130.3, 133.65, 133.74 (PhSi, benzoate), 135.4 (C(11)), 142.1 (C(12)), 167.4 (benzoate), 171.0 (4-Ac), 210.9 (C(9)) ppm. Anal. Calcd for C<sub>37</sub>H<sub>46</sub>O<sub>10</sub>Si: 1/2H<sub>2</sub>O: C, 64.61; H, 6.89. Found: C, 64.72; H, 6.81.

**[0081] 7-Dimethylisopropylsilyl-10-DAB.** To a solution of 10-DAB (97 mg, 0.18 mmol) in pyridine (1 mL) at -10 °C was added dimethylisopropylsilyl chloride (580 mL, 3.57 mmol, 20 equiv) under N<sub>2</sub>. The reaction mixture was stirred at -10 °C for 3 h. EtOAc (10 mL) was added, and the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (150 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:2) as the eluent to give 7-dimethylisopropyl-10-DAB as a white solid: yield, 107 mg (93%). mp 229-230 °C; [α]<sub>D</sub><sup>20</sup> -56° (CHCl<sub>3</sub>, c = 0.62); <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 0.05, 0.06 (2 s, 6 H, Me<sub>2</sub>Si), 0-70 (m, 1 H, CHSi), 0.90, 0.92 (2 dd, J = 7.4, 1.7, 6 H, Me<sub>2</sub>CH), 1.09 (s, 6 H, Me16, Me17), 1.56 (s, 1 H, 1-OH), 1.74 (s, 3 H, Me19), 1.89 (ddd, J = 14.4, 10.6, 2.1 Hz, 1 H, H6b), 1.99 (d, J = 5.0 Hz, 1 H, 13-OH), 2.09 (d, J = 1.4, 3 H, Me18), 2.28 (d, J = 7.9, 2 H, H14a, H14b), 2.29 (s, 3 H, 4-Ac), 2.44 (ddd, J = 14.4, 9.7, 6.7 Hz, 1 H, H6a), 3.96 (d, J = 7.3 Hz, 1 H, H3), 4.17 (d, J = 8.3 Hz, 1 H, H20b), 4.24 (d, J = 2.2 Hz, 1 H, 10-OH), 4.31 (d, J = 8.3 Hz, 1 H, H20a), 4.38 (dd, J = 10.6, 6.7 Hz, 1 H, H7), 4.85 (m, 1 H, H13), 4.95 (dd, J = 9.7, 2.1 Hz, 1 H, H5), 5.15 (d, J = 2.2 Hz, 1 H, H10), 5.61 (d, J = 7.3 Hz, 1 H, H2), 7.47 (dd, J = 8.2, 7.5 Hz, 2 H, benzoate, m), 7.60 (tt, J = 7.5, 1.4 Hz, 1 H, benzoate, p), 8.10 (d, J = 8.2, 1.4 Hz, 2 H, benzoate, o) ppm. <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>) δ -4.6, -3.3 (Me<sub>2</sub>Si), 9.7 (Me

(19)), 14.8, 14.9(CHSi, Me(18))16.4, 16.5(Me<sub>2</sub>CH), 19.4(4-Ac), 22.4, 26.7(Me16, Me17), 37.3(C(6)), 38.7(C(14)), 42.7(C(15)), 47.0(C(3)), 58.0(C(8)), 68.0(C(13)), 73.1, 74.7, 75.0(C(7), C(2), C(10), C(20)), 78.9(C(1)), 80.9(C(4)), 84.3(C(5)), 128.8, 129.8, 130.3, 133.7(benzoate), 135.7(C(11)), 142.0(C(12)), 167.4(benzoate), 171.1(4-Ac), 210.8(C(9)) ppm. Anal. Calcd for C<sub>34</sub>H<sub>48</sub>O<sub>10</sub>Si. H<sub>2</sub>O: C, 61.61; H, 7.60. Found: C, 61.30; H, 7.35.

5 **[0082]** 7-Tribenzylsilyl-10-DAB. To a mixture of 10-DAB (62 mg, 0.11 mmol), imidazole (280 mg, 4.11 mmol, 36 equiv) and tribenzylsilyl chloride (364 mg, 1.14 mmol, 10 equiv) was added DMF (0.4 mL) under N<sub>2</sub>. The reaction mixture was stirred at room temperature for 3 h. EtOAc (30 mL) was added, and the solution was filtered through a short column of silica gel. The silica gel was washed with EtOAc (150 mL), and the solution was concentrated under reduced pressure. The residue was purified twice by flash column chromatography, first time using EtOAc: hexanes (1:2) as the eluent, second time using EtOAc: CH<sub>2</sub>Cl<sub>2</sub> as the eluent, and dried overnight *in vacuo* to give the 7-tribenzylsilyl-10-DAB as a white solid: yield, 88 mg (91%). mp 161-163 °C; IR 3690, 2928, 2890, 1712, 1600 cm<sup>-1</sup>; [α]<sub>D</sub><sup>25</sup> -46° (CHCl<sub>3</sub>, c = 0.46); <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 1.10(s, 3 H, Me17, Me16), 1.56(s, 1 H, 1-OH), 1.71(ddd, J = 14.2, 10.9, 2.0 Hz, 1 H, H6b), 1.74(s, 3 H, Me19), 2.00(d, J = 5.1 Hz, 1 H, 13-OH), 2.07(ddd, J = 14.2, 9.6, 6.6 Hz, 1 H, H6a), 2.10(d, J = 1.2, 3 H, Me18), 2.12(s, 6 H, (PhCH<sub>2</sub>)<sub>3</sub>Si), 2.27(d, J = 7.5 Hz, 2 H, H14a, H14b), 2.27(s, 3 H, 4-Ac), 3.99(d, J = 7.0 Hz, 1 H, H3), 4.16(d, J = 8.5 Hz, 1 H, H20b), 4.18(d, J = 2.2 Hz, 1 H, 10-OH), 4.28(d, J = 8.5 Hz, 1 H, H20a), 4.58(dd, J = 10.9, 6.6 Hz, 1 H, H7), 4.81(dd, J = 9.6, 2.0 Hz, 1 H, H5), 4.89(m, 1 H, H13), 5.21(d, J = 2.2 Hz, 1 H, H10), 5.61(d, J = 7.0 Hz, 1 H, H2), 6.93, 7.09, 7.20(3 m, 15 H, (PhCH<sub>2</sub>)<sub>3</sub>Si), 7.48(dd, J = 8.1, 7.5 Hz, 2 H, benzoate, m), 7.61(tt, J = 7.5, 1.3 Hz, 1 H, benzoate, p), 8.10(d, J = 8.1, 1.3 Hz, 2 H, benzoate, o) ppm. <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>) δ 9.9(Me(19)), 15.0(Me(18)), 19.5(4-Ac), 22.4, 26.7(Me16, Me17), 23.6(Si(CH<sub>2</sub>Ph)<sub>3</sub>), 36.9(C(6)), 38.7(C(14)), 42.7(C(15)), 46.8(C(3)), 58.0(C(8)), 68.0(C(13)), 74.4, 74.9, 75.0(C(7), C(2), C(10), C(20)), 78.8(C(1)), 80.8(C(4)), 84.1(C(5)), 124.9, 128.7, 128.8, 129.1, 129.8, 130.3, 133.8, 137.8(Si(CH<sub>2</sub>Ph)<sub>3</sub>, benzoate), 135.5(C(11)), 142.2(C(12)), 167.4(benzoate), 170.9(4-Ac), 210.8(C(9)) ppm. Anal. Calcd for C<sub>50</sub>H<sub>56</sub>O<sub>10</sub>Si. 1/2H<sub>2</sub>O: C, 70.32; H, 6.73. Found: C, 70.11; H, 6.57.

#### Example 4

##### Selective acylation of 10-acyl-10-DAB

25 **[0083]** 10-Alloc-7-p-Nitrobenzyloxycarbonyl-10-DAB. To a mixture of 10-alloc-10-DAB (33 mg, 0.053 mmol) and DMAP (19.3 mg, 0.16 mmol, 3 equiv) in dichloromethane (4 mL) at 0 °C was added a dichloromethane solution (1 mL) of p-nitrobenzyl chloroformate (23 mL, 0.11 mmol, 2 equiv) under N<sub>2</sub>. The reaction mixture was stirred at 0 °C for 4 h. EtOAc (10 mL) was added, the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:2) as the eluent and dried overnight *in vacuo* to give 10-alloc-7-p-nitrobenzyloxycarbonyl-10-DAB as a colorless solid: yield, 34 mg (92%).

35 **[0084]** 7-Benzyl oxycarbonyl baccatin III. To a stirred solution of baccatin III (100 mg, 0.168 mmol) in methylene chloride under N<sub>2</sub> at room temperature was added 4-dimethylaminopyridine (204 mg, 1.68 mmol) followed by addition of benzyl chloroformate (240 mL, 1.68 mmol). The reaction mixture was stirred at room temperature and the progress of the reaction was monitored by TLC. After about 4 h the reaction was complete. The mixture was diluted with EtOAc (10 mL) and was transferred to a separatory funnel containing 50 mL of a 50% EtOAc/Hexanes. The mixture was washed with saturated sodium bicarbonate and the organic layer was separated. The aqueous layer was washed with 20 mL of 50% EtOAc/Hexanes. The combined organic layers were washed with brine, dried over MgSO<sub>4</sub>, and concentrated under reduced pressure. The crude product was passed through a short column to give 115 mg (95%) of a white solid m.p. 245-248 °C; [α]<sub>D</sub><sup>25</sup> -60.5° c (C=0.007, CHCl<sub>3</sub>). <sup>1</sup>H NMR (CDCl<sub>3</sub>, 400 MHz) δ 8.10(d, J=9.6 Hz, 2H, o-benzoate), 7.60-6.8(m, 8H, benzoate, Bn), 6.45(s, 1H, H10), 5.63(d, J=6.9 Hz, 1H, H2b), 5.56(dd, J=10.6, 7.2 Hz, 1H, H7), 5.56(dd, J=18.5, 12.0 Hz, 2H, Bn), 4.97(d, J=10.6, 1H, H5), 4.87(m, 1H, H13), 4.31(d, J=10.5, 1H, H20a), 4.15(d, J=10.5, 1H, H20b), 4.02(d, J=6.9, 1H, H3), 2.61(m, 1H, H6a), 2.30(m, 2H, H14's), 2.29(s, 3H, 4Ac), 2.18(s, 3H, 10Ac), 2.15(br s, 3H, Me18), 2.08(d, J=5.2 Hz, 13OH), 1.94(m, 1H, 6b), 1.79(s, 3H, Me19), 1.58(s, 1H, 10H), 1.14(s, 3H, Me16), 1.09(s, 3H, Me17).

40 **[0085]** 7-Allyloxycarbonyl baccatin III. To a stirred solution of baccatin III (30 mg, 0.051 mmol) in methylene chloride (1 mL) under N<sub>2</sub> at room temperature, was added 4-dimethylaminopyridine (62.3 mg, 0.51 mmol) followed by addition of allyl chloroformate (54 mL, 0.51 mmol). The reaction mixture was stirred at room temperature and the progress of the reaction was followed by TLC. After about 1.5 h the reaction was complete. The mixture was diluted by EtOAc (5 mL) and was transferred to a separatory funnel containing 50 mL of a 50% EtOAc/Hexanes. The mixture was washed with saturated sodium bicarbonate and the organic layer was separated. The aqueous layer was washed with 10 mL of 50% EtOAc/Hexanes, the combined organic layers were washed with brine, dried over MgSO<sub>4</sub>, and concentrated under reduced pressure. The crude product was passed through a short column to give 33.1 mg (97%) of a white solid m.p. 239-244 °C; [α]<sub>D</sub><sup>25</sup> -61.5° c (0.01, CHCl<sub>3</sub>). <sup>1</sup>H NMR (CDCl<sub>3</sub>, 500 MHz) δ 8.12(d, J=8.3 Hz, 2H, o-benzoate), 7.66-7.45(m, 3H, benzoate), 6.43(s, 1H, H10), 5.97(m, 1H, int. allyl), 5.64(d, J=7.0 Hz, 1H, H2b), 5.54(dd, J=10.5, 7.0 Hz, 1H, H7), 5.28(m, 2H, ext. allyl), 4.97(d, J=9.6 Hz, 1H, H5), 4.87(m, 1H, H13), 4.67(m, 2H, CH<sub>2</sub>allyl), 4.31(d, J=8.5 Hz, 1H, H20a),

4.17(d, J=8.5, 1H, H20b), 4.02(d, J=7.0, 1H, H3), 2.64(m, 1H, H6a), 2.30(d, J=8.0 Hz, 2H, H14's), 2.29(s, 3H, 4Ac), 2.16(s, 3H, 10Ac), 2.15(br s, 3H, Me18), 2.01(d, J=5 Hz, 130H), 1.96(m, 1H, 6b), 1.81(s, 3H, Me19), 1.58(s, 1H, 10H), 1.15(s, 3H, Me16), 1.02(s, 3H, Me17).

## 5 Example 5

### Selective ketalization of 10-acyl-10-DAB

[0086] *7-MOP baccatin III*. To a solution of baccatin III (101 mg, 0.172 mmol) in THF (8 mL) at -20 °C was added 2-methoxypropene (0.66 mL, 6.89 mmol, 40 equiv), followed by the addition of a catalytic amount of toluenesulfonic acid (0.1 M solution in THF, 43  $\mu$ L, 0.004 mmol, 0.025 equiv) under N<sub>2</sub>. The reaction mixture was stirred at -20 °C for 3 h. TLC analysis indicated complete consumption of the starting material and the formation of desired product as the only major product. Triethylamine (0.5 mL) was added, and the solution was warmed to room temperature, diluted with EtOAc (100 mL), washed with saturated aqueous NaHCO<sub>3</sub> solution, dried over Na<sub>2</sub>SO<sub>4</sub>, and concentrated under reduced pressure. The residue was dried in *vacuo* overnight to give 112 mg (99%) of crude product. Recrystallization of the crude product from EtOAc/hexanes gave 105 mg (93%) of 7-MOP baccatin III as a white crystal, mp 181-183 °C; <sup>1</sup>H NMR (500 MHz, C<sub>6</sub>D<sub>6</sub>)  $\delta$  1.01 (s, 3 H, Me17), 1.11(br s, 1 H, 13-OH), 1.28(s, 3 H, Me16), 1.39, 1.78(2 s, 6 H, Me<sub>2</sub>CO), 1.62 (s, 1 H, 1-OH), 1.78(s, 3 H, 10-Ac), 1.92(s, 3 H, 4-Ac), 2.09(s, 3 H, Me18), 2.12(s, 3 H, Me19), 2.14(ddd, J = 15.0, 10.9, 2.2 Hz, 1 H, H6b), 2.18(dd, J = 15.6, 9.4 Hz, 1 H, H14b), 2.31(dd, J = 15.6, 7.0 Hz, 1 H, H14b), 2.97(s, 3 H, MeO), 3.15 (ddd, J = 15.0, 9.9, 6.7 Hz, 1 H, H6a), 4.08(d, J = 7.0 Hz, 1 H, H3), 4.24(m, 1 H, H13), 4.33(d, J = 8.3 Hz, 1 H, H20b), 4.41(d, J = 8.3 Hz, 1 H, H20a), 4.78(dd, J = 10.9, 6.7 Hz, 1 H, H7), 4.97(dd, J = 9.9, 2.2 Hz, 1 H, H5), 5.95(d, J = 7.0 Hz, 1 H, H2), 6.79(s, 1 H, H10), 7.15(m, 3 H, benzoate, m, p), 8.28(d, J = 8.0 Hz, 2 H, benzoate, o) ppm; Anal. Calcd for C<sub>35</sub>H<sub>46</sub>O<sub>12</sub>: C, 63.82; H, 7.04. Found: C, 63.72; H, 7.07.

## 25 Example 6

### Example 7

### Selective silylation of 10-acyl-10-DAB

[0087] *7-dimethylisopropylsilyl baccatin III*. To a stirred solution of baccatin III (30 mg, 0.051 mmol) in pyridine (0.6 mL) at 0°C under N<sub>2</sub>, was added chlorodimethyl-isopropylsilane (160  $\mu$ L, 1.02 mmol). The reaction mixture was stirred at that temperature and the progress of the reaction was monitored by TALC. After about 1.5 h the reaction was complete. Ethyl acetate (5 mL) was added and the solution was transferred to a separatory funnel containing 50 mL of a 50% EtOAc/Hexanes. The mixture was washed with saturated sodium bicarbonate and the organic layer was separated. The aqueous layer was extracted with 10 mL of 50% EtOAc/Hexanes and the combined organic layers were washed with saturated sodium chloride, dried over MgSO<sub>4</sub>, concentrated under reduced pressure. The crude product was passed through a short silica gel column to give 33.9 mg (97%) of a white solid m.p. 204-207 °C; [ $\alpha$ ]<sub>D</sub><sup>25</sup> -58.6° c (0.009, CHCl<sub>3</sub>). <sup>1</sup>H NMR (CDCl<sub>3</sub>, 500MHz),  $\delta$  8.10(d, J=8.4Hz, 2H, o-benzoate), 7.60-7.20 (m, 3H, benzoate), 6.4 (s, 1H, H10), 5.64 (d, J=7.1Hz, 1H, H2b), 4.95 (d, J=4.9Hz, 1H, H5), 4.84 (m, 1H, H13), 4.44 (dd, J=10.4, 6.8 Hz, 1H, H7), 4.30(d, J=8.3 Hz, 1H, H20a), 4.14 (d, J=8.3 Hz, 1H, H20b), 4.15(d, J=7.2 Hz, 1H, H3), 2.49(m, 1H, H6a), 2.23 (m, 2H, H14's), 2.28 (s, 3H, 4Ac), 2.18 (br s, 3H, Me 18), 2.17 (s, 3H, 10Ac), 2.01(d, J=5.0 Hz, 13 OH), 1.86(m, 1H, 6b), 1.69(s, 3H, Me19), 1.61(s, 1H, 10H), 1.20(s, 3H, Me16), 1.05 (s, 3H, Me17), 0.87 (d, J=7.1 Hz, 6H, i-pr), 0.73(m, 1H, i-pr), 0.09(s, 6H, Me2Si).

[0088] *7-dimethylphenylsilyl baccatin III*. To a stirred solution of baccatin III (20 mg, 0.034 mmol) in THF (1.25 mL) at -10°C under N<sub>2</sub>, was added chlorodimethylphenyl-silane (68  $\mu$ L, 0.41 mmol), followed by addition of pyridine (250 mL, 3.1 mmol). The reaction mixture was stirred at that temperature and the progress of the reaction was monitored by TLC. After about one hour the reaction was complete. Ethyl acetate (5 mL) was added and the solution was transferred to a separatory funnel containing 30 mL of 50% EtOAc/Hexanes. The mixture was washed with saturated sodium bicarbonate and the organic layer was separated. The aqueous layer was extracted with 10 mL of 50% EtOAc/Hexanes and the combined organic layers were washed with saturated sodium chloride, dried over MgSO<sub>4</sub>, concentrated under reduced pressure. The crude product was passed through a short silica gel column to give 24.1 mg (98%) of a white solid m.p. 210-213°C; [ $\alpha$ ]<sub>D</sub><sup>25</sup> -58.3.5° c (0.005, CHCl<sub>3</sub>). <sup>1</sup>H NMR (CDCl<sub>3</sub>, 500 MHz)  $\delta$  8.35(d, J=8.5 Hz, 2H, o-benzoate), 7.627.25 (m, 8H, benzoate, phenyl), 6.42 (s, 1H, H10), 5.64 (d, J=6.9Hz, 1H, H2b), 4.84 (m, 1H, H5), 4.81 (m, 1H, H13), 4.46 (dd, J=10.6, 6.9 Hz, 1H, H7), 4.21 (d, J=8.5 Hz, 1H, H20a), 4.14 (d, J=8.5 Hz, 1H, H20b), 3.85(d, J=6.9 Hz, 1H, H3), 2.34(m, 1H, H6a), 2.26 (d, J=8 Hz, 2H, H14's), 2.24 (s, 3H, 4Ac), 2.15(s, 3H, 10Ac), 2.02(br d, J=1 Hz, 3H, Me 18), 1.93 (d, J=5 Hz, 1H, 130H), 1.77(m, 1H, 6b), 1.72(s, 3H, Me19), 1.59 (s, 1H, 10H), 1.20 (s, 3H, Me16), 1.05(s, 3H, Me17), 0.446(s, 3H, Me Si), 0.335(s, 3H, Me Si).

[0089] *7-dimethylphenylsilyl-10-propionyl-10-DAB*. To a stirred solution of 10-propionyl-10-DAB (0.200 g, 0.333 mmol)

in THF (12 mL) at -10°C, was added chlorodimethyl-phenylsilane (0.668 mL, 4.00 mmol) followed by pyridine dropwise (2.48 mL, 30.64 mmol). The reaction was stirred for 90 minutes. Ethyl acetate (20 mL) was added and the solution transferred to a separatory funnel containing 100 mL of 50% EtOAc/Hexanes. The mixture was washed with saturated sodium bicarbonate and the organic layer separated. The aqueous layer was extracted with 50% EtOAc/Hexanes (30 mL) and the combined organic extracts washed with saturated sodium chloride, dried over Na<sub>2</sub>SO<sub>4</sub>, concentrated *in vacuo*. The crude solid was then purified with flash column chromatography using 50% EtOAc/hexane as eluent to give 7-dimethylphenylsilyl-10-propionyl-10-DAB (0.242 g, 99%) as a solid. <sup>1</sup>H NMR (CDCl<sub>3</sub>, 500 MHz), δ 0.34, 0.45(2 s, 6 H, Me<sub>2</sub>Si), 1.05 (s, 3 H, Me17), 1.20 (t, J = 7.5 Hz, 3 H, CH<sub>3</sub>CH<sub>2</sub>), 1.21 (s, 3 H, Me16), 1.60(s, 1 H, 1-OH), 1.72(s, 3 H, Me19), 1.78(ddd, J = 14.5, 10.0, 2.0 Hz, 1 H, H6b), 2.04(m, 1 H, 13-OH), 2.05 (s, 3 H, Me18), 2.27 (m, 2 H, H14a, H14b), 2.25 (s, 3 H, 4-Ac), 2.34(ddd, J = 14.5, 9.5, 7.0 Hz, 1 H, H6a), 2.42, 2.49 (2 dq, J = 16.5, 7.5 Hz, 6 H, CH<sub>3</sub>CH<sub>2</sub>), 3.87 (d, J = 7.5 Hz, 1 H, H3), 4.14 (d, J = 8.0 Hz, 1 H, H20b), 4.27(d, J = 8.0 Hz, 1 H, H20a), 4.47(dd, J = 10.0, 7.0 Hz, 1 H, H7), 4.82(m, 1 H, H13), 4.85(dd, J = 9.5, 2.0 Hz, 1 H, H5), 5.64(d, J = 7.5 Hz, 1 H, H2), 6.44(s, 1 H, H10), 7.32-7.36, 7.55-7.57 (2 m, 5 H, PhSi), 7.46(m, 2 H, benzoate, m), 7.59(m, 1 H, benzoate, p), 8.10 (d, J = 8.0 Hz, 2 H, benzoate, o) ppm.

[0090] 7-Dimethylphenylsilyl-10-cyclopropanecarbonyl-10-DAB. To a solution of 10-cyclopropanecarbonyl-10-DAB (680 mg, 1.1 mmol) in THF (25 mL) were added with stirring pyridine (3.5 mL) and then chlorodimethyl-phenylsilane (1.8 mL, 11 mmol) at -10 °C under N<sub>2</sub>. The solution was stirred till the reaction completed. Then quenched with sat. NaHCO<sub>3</sub> (20 mL). The mixture was extracted with EtOAc (2 x 250 mL). The combined organic layers were washed with brine (2 x 10 mL), dried and filtered. Concentration of the filtrate *in vacuo* and followed by flash chromatography (hexane: EtOAc, 4:1) gave 7-Dimethyl-phenylsilyl-10-cyclopropane-carbonyl-10-DAB (816 mg, -100%). <sup>1</sup>H NMR (CDCl<sub>3</sub>, 500 MHz), δ 0.32, 0.43(2 s, 6 H, Me<sub>2</sub>Si), 0.91, 1.00, 1.17(3 m, 5 H, cyclopropyl), 1.07(s, 3 H, Me17), 1.21(s, 3 H, Me16), 1.73(s, 3 H, Me19), 1.74(s, 1H, 1-OH), 1.78(ddd, J = 14.4, 10.5, 2.1 Hz, 1 H, H6b), 2.04(m, 1 H, 13-OH), 2.05(d, J = 1.5 Hz, 3 H, Me18), 2.24 (s, 3 H, 4-Ac), 2.26 (m, 2 H, H14a, H14b), 2.34 (ddd, J = 14.4, 9.5, 6.7 Hz, 1 H, H6a), 3.87(d, J = 7.0 Hz, 1 H, H3), 4.15(d, J = 8.2 Hz, 1 H, H20b), 4.26(d, J = 8.2 Hz, 1 H, H20a), 4.46(dd, J = 10.5, 6.7 Hz, 1 H, H7), 4.82(m, 1 H, H13), 4.85(dd, J = 9.5, 2.1 Hz, 1 H, H5), 5.65 (d, J = 7.0 Hz, 1 H, H2), 6.44(s, 1 H, H10), 7.32-7.36, 7.55-7.57(2 m, 5 H, PhSi), 7.46(m, 2 H, benzoate, m), 7.59(m, 1 H, benzoate, p), 8.10(d, J = 8.0 Hz, 2 H, benzoate, o) ppm.

### Example 8

[0092] 7-p-Nitrobenzyloxycarbonyl-10-DAB. To a THF solution (1 mL) of 10-alloc-7-p-nitrobenzyloxycarbonyl-10-DAB (34 mg, 0.048 mmol) at room temperature was added a THF solution (1 mL) of formic acid (19 mL, 0.48 mmol, 10 equiv) and butylamine (47 mL, 0.48 mmol, 10 equiv), followed by the addition of Pd(PPh<sub>3</sub>) under N<sub>2</sub>. The reaction mixture was stirred at room temperature for 0.5 h. EtOAc (10 mL) was added, and the solution was quickly filtered through a short column of silica gel. The silica gel was washed with EtOAc (100 mL), and the solution was concentrated under reduced pressure. The residue was purified by flash column chromatography using EtOAc: hexanes (1:2) as the eluent and dried *in vacuo* to give 7-p-nitrobenzyloxycarbonyl-10-DAB as a colorless solid: yield, 28 mg (93%). [α]<sub>D</sub><sup>20</sup> -38° (CHCl<sub>3</sub>, c = 0.48); <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ 1.06(s, 3 H, Me16), 1.09(s, 3 H, Me17), 1.55(s, 1H, 1-OH), 1.86(s, 3 H, Me19), 2.01 (ddd, J = 14.4, 10.7, 2.0 Hz, 1 H, H6b), 2.03(d, J = 5.1 Hz, 1 H, 13-OH), 2.09(d, J = 1.3, 3 H, Me18), 2.28(m, 2 H, H14a, H14b), 2.30(s, 3 H, 4-Ac), 2.62(ddd, J = 14.4, 9.5, 7.3 Hz, 1 H, H6a), 3.89(d, J = 2.0 Hz, 1 H, 10-OH), 4.08(d, J = 6.9 Hz, 1 H, H3), 4.20(d, J = 8.4 Hz, 1 H, H20b), 4.34(d, J = 8.4 Hz, 1 H, H20a), 4.88(m, 1 H, H13), 4.96(dd, J = 9.5, 2.0 Hz, 1 H, H5), 5.19(d, J = 13.3, 1 H, CHH'OC(O)), 5.26 (d, J = 13.3, 1 H, CHH'OC(O)), 5.36 (dd, J = 10.7, 7.3 Hz, 1 H, H7), 5.40 (d, J = 2.0 Hz, 1 H, H10), 5.64(d, J = 6.9 Hz, 1 H, H2), 7.48(dd, J = 8.1, 7.5 Hz, 2 H, benzoate, m), 7.52(d, J = 8.7, 2 H, NO<sub>2</sub>C<sub>6</sub>H<sub>4</sub>), 7.61 (tt, J = 7.5, 1.3 Hz, 1 H, benzoate, p), 8.10(d, J = 8.1, 1.3 Hz, 2 H, benzoate, o), 8.26(d, J = 8.7, 2 H, NO<sub>2</sub>C<sub>6</sub>H<sub>4</sub>) ppm. <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>) δ 10.5(Me(19)), 14.6(Me(18)), 19.4(4-Ac), 22.2, 26.4(Me16, Me17), 33.2(C(6)), 38.7(C(14)), 42.4(C(15)), 46.5(C(3)), 56.5(C(8)), 67.9, 68.3 (C(13), OCH<sub>2</sub>Ph-NO<sub>2</sub>-P), 74.7, 75.2, 76.8(C(7), C(2), C(10), C(20)), 78.8(C(1)), 80.4(C(4)), 83.6(C(5)), 124.1, 128.4, 128.9, 130.3, 133.9 (OCH<sub>2</sub>Ph-NO<sub>2</sub>-p, benzoate), 135.0(C(11)), 142.4, 143.0 (OCH<sub>2</sub>Ph-NO<sub>2</sub>-p, C(12)), 154.2 (OC(O)O), 167.3 (benzoate), 171.1(4-Ac), 211.6(C(9))ppm.

### Example 9

[0093] Selective Esterification of the C-10 Hydroxyl of 10-DAB using the catalytic DyCl<sub>3</sub> reaction: A solution of butyric anhydride (0.55 mmol) in THF (1.32 ml) was added, under a nitrogen atmosphere, to a solid mixture of 10-DAB (30 mg, 0.055 mmol) and DyCl<sub>3</sub> (1.3 mg, 10 mol.% wrt 10-DAB). The resulting suspension was stirred at room temperature until judged complete by TLC (2:1 EtOAc/Hexane). The reaction was diluted with EtOAc and washed three times with saturated NaHCO<sub>3</sub> solution. The combined bicarbonate washings were extracted three times with EtOAc, these combined organics were dried (Na<sub>2</sub>SO<sub>4</sub>) and the solvent evaporated. The crude product was triturated with hexanes and the mother liquors decanted away. Crystallization from EtOAc/hexanes yielded 10-butyl-10-DAB identical to that isolated from the CeCl<sub>3</sub> catalysed reaction.



## Example 10

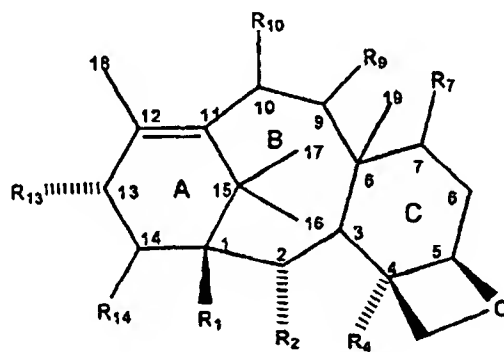
[0094] *Selective Esterification of the C-10 Hydroxyl of 10-DAB using the catalytic  $\text{YbCl}_3$  reaction:* A solution of butyric anhydride (0.55 mmol) in THF (1.32 ml) was added, under a nitrogen atmosphere, to a solid mixture of 10-DAB (30 mg, 0.055 mmol) and  $\text{YbCl}_3$  (1.3 mg, 10 mol.% wrt 10-DAB). The resulting suspension was stirred at room temperature until judged complete by TLC (2:1 EtOAc/Hexane). The reaction was diluted with EtOAc and washed three times with saturated  $\text{NaHCO}_3$  solution. The combined bicarbonate washings were extracted three times with EtOAc, these combined organics were dried ( $\text{Na}_2\text{SO}_4$ ) and the solvent evaporated. The crude product was triturated with hexanes and the mother liquors decanted away. Crystallization from EtOAc/hexanes yielded 10-butyryl-10-DAB identical to that isolated from the  $\text{CeCl}_3$  catalysed reaction.

[0095] In view of the above, it will be seen that the several objects of the invention are achieved.

[0096] As various changes could be made in the above compositions without departing from the scope of the invention, it is intended that all matter contained in the above description be interpreted as illustrative and not in a limiting sense.

## Claims

1. A process for the acylation of a C(10) hydroxy group of a taxane, the process comprising treating the taxane with an acylating agent selected from dicarbonates, thiodicarbonates, and isocyanates in a reaction mixture containing less than one equivalent of a base for each equivalent of taxane to form a C(10) acylated taxane.
2. A process according to claim 1, wherein the taxane has C(7) and C(10) hydroxy groups and the acylating agent selectively reacts with the C(10) hydroxy group.
3. A process according to claim 1 or claim 2, wherein the taxane reacted with the acylating agent is 10-deacetyl baccatin III.
4. A process according to any one of claims 1 to 3, wherein the taxane has the structure:



wherein

$\text{R}_1$  is hydrogen, hydroxy, protected hydroxy, or together with  $\text{R}_{14}$  or  $\text{R}_2$  forms a carbonate;

$\text{R}_2$  is keto,  $-\text{OT}_2$ , acyloxy, or together with  $\text{R}_1$  forms a carbonate;

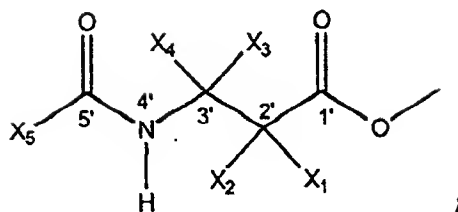
$\text{R}_4$  is  $-\text{OT}_4$ , or acyloxy;

$\text{R}_7$  is hydrogen, halogen,  $-\text{OT}_7$ ,  $-\text{OCOZ}_7$ , or  $-\text{OCOOZ}_7$ ;

$\text{R}_9$  is hydrogen, keto,  $-\text{OT}_9$ ,  $-\text{OCOZ}_7$ , or  $-\text{OCOOZ}_7$ ;

$\text{R}_{10}$  is hydroxy;

$\text{R}_{13}$  is hydroxy, protected hydroxy, keto, or



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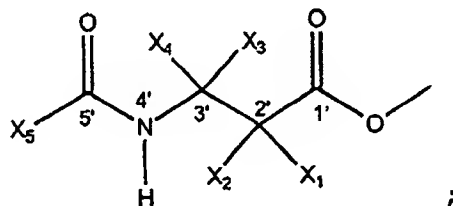
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- $R_{14}$  is hydrogen,  $-OT_{14}$ , acyloxy, or together with  $R_1$  forms a carbonate;  
 $T_2, T_4, T_7, T_9$ , and  $T_{14}$  are independently hydrogen or hydroxy protecting group;  
 $X_1$  is  $-OX_6$ ,  $-SX_7$ , or  $-NX_8X_9$ ;  
 $X_2$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;  
 $X_3$  and  $X_4$  are independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;  
 $X_5$  is  $-X_{10}$ ,  $-OX_{10}$ ,  $-SX_{10}$ ,  $-NX_8X_{10}$ , or  $-SO_2X_{11}$ ;  
 $X_6$  is hydrocarbyl, substituted hydrocarbyl, heteroaryl, hydroxy protecting group or a functional group which increases the water solubility of the taxane derivative;  
 $X_7$  is hydrocarbyl, substituted hydrocarbyl, heteroaryl, or sulfhydryl protecting group;  
 $X_8$  is hydrogen, hydrocarbyl, or substituted hydrocarbyl;  
 $X_9$  is an amino protecting group;  
 $X_{10}$  is hydrocarbyl, substituted hydrocarbyl, or heteroaryl;  
 $X_{11}$  is hydrocarbyl, substituted hydrocarbyl, heteroaryl,  $-OX_{10}$ , or  $-NX_8X_{14}$ ; and  
 $X_{14}$  is hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

5. A process according to claim 4, wherein  
 $R_1$  is hydroxy or together with  $R_{14}$  or  $R_2$  forms a carbonate;  
 $R_2$  is  $-OCOZ_2$ ,  $-OCOOZ_2$ , or together with  $R_1$  forms a carbonate;  
 $R_4$  is  $-OCOZ_4$ ;  
 $R_9$  is hydrogen or keto;  
 $R_{13}$  is hydroxy, protected hydroxy, or

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- $R_{14}$  is hydrogen, hydroxy, protected hydroxy, or together with  $R_1$  forms a carbonate;  
 $X_1$  is  $-OX_6$  or  $-NX_8X_9$ ;  
 $X_2$  is hydrogen, hydrocarbyl, or substituted hydrocarbyl;  
 $X_3$  and  $X_4$  are independently hydrogen, hydrocarbyl, substituted hydrocarbyl, or heteroaryl;  
 $X_5$  is  $-X_{10}$ ,  $-OX_{10}$ , or  $-NX_8X_{10}$ ;  
 $X_6$  is a hydroxy protecting group;  
 $X_8$  is hydrogen, hydrocarbyl, or substituted hydrocarbyl;  
 $X_9$  is an amino protecting group;  
 $X_{10}$  is hydrocarbyl, substituted hydrocarbyl, or heteroaryl; and  
 $Z_2$  and  $Z_4$  are independently hydrocarbyl, substituted hydrocarbyl, or heteroaryl.

6. A process according to any one of claims 1 to 5, wherein the reaction mixture contains a Lewis acid.

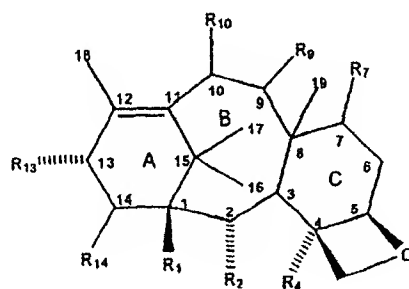
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7. A process according to claim 6, wherein the Lewis acid is selected from the halides or triflates of the Group IB, IIB, IIIB, IVB, VB, VIIB, VIII, IIIA, IVA, lanthanide and actinide elements.

8. A process according to claim 7, wherein the Lewis acid is selected from zinc chloride, stannic chloride, cerium trichloride, cuprous chloride, lanthanum trichloride, dysprosium trichloride and ytterbium trichloride.
9. A process according to any one of claims 1 to 8,  
wherein the C(10) acylated taxane comprises a C(7) hydroxy group and the process additionally comprises treating the C(10) acylated taxane with a silylating agent to silylate the C(7) hydroxy group.
10. A process according to any one of claims 1 to 8,  
wherein the C(10) acylated taxane comprises a C(7) hydroxy group and the process additionally comprises treating the C(10) acylated taxane with an acylating agent to acylate the C(7) hydroxy group.
11. A process according to claim 9 or claim 10, wherein the C(10) acylated taxane is baccatin III.
12. A process according to any one of claims 1 to 8,  
wherein the C(10) acylated taxane comprises a C(13) hydroxy, metallic oxide, or ammonium oxide substituent and the process additionally comprises the step of esterifying the C(10) acylated taxane by treating the C(10) acylated taxane with a side chain precursor selected from  $\beta$ -lactams, oxazolines, oxazolidine carboxylic acids, oxazolidine carboxylic acid anhydrides, and isoserine derivatives.

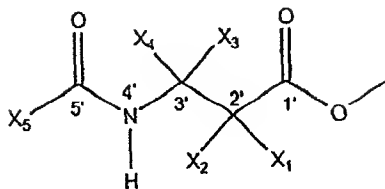
## Patentansprüche

1. Verfahren zur Acylierung einer Hydroxygruppe an C(10) eines Taxans, bei dem man das Taxan mit einem unter Dicarbonaten, Thiodicarbonaten und Isocyanaten ausgewählten Acylierungsmittel in einem weniger als ein Äquivalent einer Base für jedes Äquivalent Taxan enthaltenden Reaktionsgemisch behandelt, um ein an C(10) acyliertes Taxan zu bilden.
2. Verfahren nach Anspruch 1, bei dem das Taxan Hydroxygruppen an C(7) und C(10) hat und das Acylierungsmittel selektiv mit der Hydroxygruppe an C(10) reagiert.
3. Verfahren nach Anspruch 1 oder Anspruch 2, bei dem das mit dem Acylierungsmittel umgesetzte Taxan 10-Deacetylbaccatin III ist.
4. Verfahren nach einem der Ansprüche 1 bis 3, bei dem das Taxan die Struktur



hat, worin

- $R_1$  Wasserstoff, Hydroxy, geschütztes Hydroxy ist oder zusammen mit  $R_{14}$  oder  $R_2$  ein Carbonat bildet,  
 $R_2$  Keto,  $-OT_2$ , Acyloxy ist oder zusammen mit  $R_1$  ein Carbonat bildet,  
 $R_4$   $-OT_4$  oder Acyloxy ist,  
 $R_7$  Wasserstoff, Halogen,  $-OT_7$ ,  $-OCOZ_7$  oder  $-OCOOZ_7$  ist,  
 $R_9$  Wasserstoff, Keto,  $-OT_9$ ,  $-OCOZ_9$  oder  $-OCOOZ_9$  ist,  
 $R_{10}$  Hydroxy ist,  
 $R_{13}$  Hydroxy, geschütztes Hydroxy, Keto oder



ist,

$R_{14}$  Wasserstoff, -QT<sub>14</sub>, Acyloxy ist oder zusammen mit  $R_1$  ein Carbonat bildet,

$T_2$ ,  $T_4$ ,  $T_7$ ,  $T_9$  und  $T_{14}$  unabhängig Wasserstoff oder eine Hydroxy-Schutzgruppe sind,

$X_1$  -OX<sub>6</sub>, -SX<sub>7</sub> oder -NX<sub>8</sub> X<sub>9</sub> ist,

$X_2$  Wasserstoff, Hydrocarbyl, substituiertes Hydrocarbyl oder Heteroaryl ist,

$X_3$  und  $X_4$  unabhängig Wasserstoff, Hydrocarbyl, substituiertes Hydrocarbyl oder Heteroaryl sind,

$X_5$  -X<sub>10</sub>, -OX<sub>10</sub>, -SX<sub>10</sub>, -NX<sub>8</sub>X<sub>10</sub> oder -SO<sub>2</sub>X<sub>11</sub> ist,

$X_6$  Hydrocarbyl, substituiertes Hydrocarbyl, Heteroaryl, eine Hydroxy-Schutzgruppe oder eine funktionelle Gruppe ist, die die Wasserlöslichkeit des Taxanderivats erhöht,

$X_7$  Hydrocarbyl, substituiertes Hydrocarbyl, Heteroaryl oder eine Sulfhydryl-Schutzgruppe ist,

$X_8$  Wasserstoff, Hydrocarbyl oder substituiertes Hydrocarbyl ist,

$X_9$  eine Amino-Schutzgruppe ist,

$X_{10}$  Hydrocarbyl, substituiertes Hydrocarbyl oder Heteroaryl ist,

$X_{11}$  Hydrocarbyl, substituiertes Hydrocarbyl, Heteroaryl, -OX<sub>10</sub>, oder -NX<sub>8</sub>X<sub>14</sub> ist, und

$X_{14}$  Wasserstoff, Hydrocarbyl, substituiertes Hydrocarbyl oder Heteroaryl ist.

#### 5. Verfahren nach Anspruch 4, bei dem

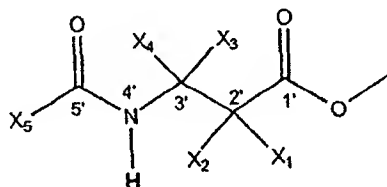
$R_1$  Hydroxy ist oder zusammen mit  $R_{14}$  oder  $R_2$  ein Carbonat bildet,

$R_2$  -OCOZ<sub>2</sub>, -OCOOZ<sub>2</sub> ist oder zusammen mit  $R_1$  ein Carbonat bildet,

$R_4$  -OCOZ<sub>4</sub> ist,

$R_9$  Wasserstoff oder Keto ist,

$R_{13}$  Hydroxy, geschütztes Hydroxy oder ist,



$R_{14}$  Wasserstoff, Hydroxy, geschütztes Hydroxy ist oder zusammen mit  $R_1$  ein Carbonat bildet,

$X_1$  -OX<sub>6</sub> oder -NX<sub>8</sub>X<sub>9</sub> ist,

$X_2$  Wasserstoff, Hydrocarbyl oder substituiertes Hydrocarbyl ist,

$X_3$  und  $X_4$  unabhängig Wasserstoff, Hydrocarbyl, substituiertes Hydrocarbyl oder Heteroaryl sind,

$X_5$  -X<sub>10</sub>, -OX<sub>10</sub> oder -NX<sub>8</sub> X<sub>10</sub> ist,

$X_6$  eine Hydroxy-Schutzgruppe ist,

$X_8$  Wasserstoff, Hydrocarbyl oder substituiertes Hydrocarbyl ist,

$X_9$  eine Amino-Schutzgruppe ist,

$X_{10}$  Hydrocarbyl, substituiertes Hydrocarbyl oder Heteroaryl ist, und

$Z_2$  und  $Z_4$  unabhängig Hydrocarbyl, substituiertes Hydrocarbyl oder Heteroaryl sind.

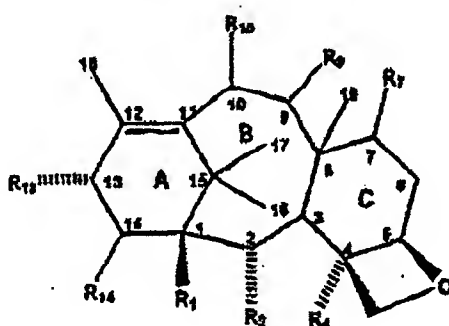
#### 6. Verfahren nach einem der Ansprüche 1 bis 5, bei dem das Reaktionsgemisch eine Lewis-Säure enthält.

#### 7. Verfahren nach Anspruch 6, bei dem die Lewis-Säure unter den Halogeniden oder Triflats der Gruppe IB, IIB, IIIB, IVB, VB, VIIB, VIIIB, IIIA, IVA, der Lanthaniden und Actinidenelemente ausgewählt ist.

8. Verfahren nach Anspruch 7, bei dem die Lewis-Säure unter Zinkchlorid, Zinn(IV)-chlorid, Certrichlorid, Kupfer(I)-chlorid, Lanthantrichlorid, Dysprosiumtrichlorid und Ytterbiumtrichlorid ausgewählt ist.
9. Verfahren nach einem der Ansprüche 1 bis 8, bei dem das an C(10) acylierte Taxan eine Hydroxygruppe an C(7) hat und das Verfahren ferner die Behandlung des an C(10) acylierten Taxans mit einem Silylierungsmittel umfaßt, um die Hydroxygruppe an C(7) zu silylieren.
10. Verfahren nach einem der Ansprüche 1 bis 8, bei dem das an C(10) acylierte Taxan eine Hydroxygruppe an C(7) hat und das Verfahren ferner die Behandlung des an C(10) acylierten Taxans mit einem Acylierungsmittel umfaßt, um die Hydroxygruppe an C(7) zu acylieren.
11. Verfahren nach Anspruch 9 oder Anspruch 10, bei dem das an C(10) acylierte Taxan Baccatin III ist.
12. Verfahren nach einem der Ansprüche 1 bis 8, bei dem das an C(10) acylierte Taxan an C(13) einen Hydroxy-, Metalloxyd- oder Ammoniumoxyd-Substituenten aufweist und das Verfahren ferner die Stufe der Veresterung des an C(10) acylierten Taxans durch dessen Behandlung mit einem Seitenkettenvorprodukt umfaßt, das unter  $\beta$ -Lactamen, Oxazolinen, Oxazolidincarbonsäuren, Oxazolidincarbonsäureanhydriden und Isoserinderivaten ausgewählt ist.

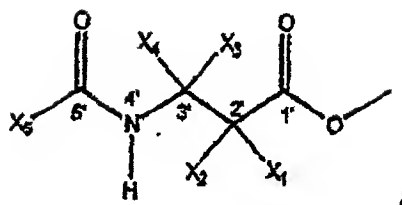
## Revendications

1. Procédé pour l'acylation d'un groupe hydroxy en C(10) d'un taxane, le procédé comprenant le traitement du taxane avec un agent acylant choisi parmi les dicarbonates, les et les isocyanates dans un mélange réactionnel contenant moins d'équivalent d'une base par de taxane pour former un taxane acylé en C(10).
2. Procédé selon la revendication 1 où le taxane a des groupes hydroxy en C(7) et C(10) et l'agent acylant réagit sélectivement avec le groupe hydroxy en C(10).
3. Procédé selon la revendication 1 ou la revendication 2 où le taxane mis à réagir avec l'agent acylant est la 10-désacétyl baccatine III.
4. Procédé selon l'une quelconque des revendications 1 à 3 où le taxane a structure :



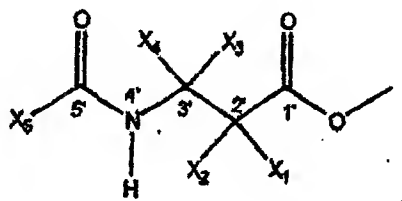
où

- $R_1$  est l'hydrogène, hydroxy, hydroxy ou forme un carbonate avec  $R_{14}$  ou  $R_2$  ;  
 $R_2$  est céto,  $-OT_2$ , acyloxy, ou forme un carbonate avec  $R_1$  ;  
 $R_4$  est  $-OT_4$  ou acyloxy ;  
 $R_7$  est l'hydrogène, un halogène,  $-OT_7$ ,  $-OCOZ_7$  ou  $-OCOOZ_7$  ;  
 $R_9$  est l'hydrogène, céto,  $-OT_9$ ,  $-OCOZ_7$  ou  $-OCOOZ_7$  ;  
 $R_{10}$  est hydroxy ;  
 $R_{13}$  est hydroxy, hydroxy protégé, céto ou



- $R_{14}$  est l'hydrogène,  $-OT_{14}$ , acyloxy, ou forme un carbonate avec  $R_1$  ;  
 $T_2, T_4, T_7, T_9$  et  $T_{14}$  sont indépendamment l'hydrogène ou un groupe protecteur d'hydroxy ;  
 $X_1$  est  $-OX_6$ ,  $-SX_7$  ou  $-NX_8X_9$  ;  
 $X_2$  est l'hydrogène, hydrocarbyle, hydrocarbyle substitué ou hétéroaryle ;  
 $X_3$  et  $X_4$  sont indépendamment l'hydrogène, hydrocarbyle, hydrocarbyle substitué ou hétéroaryle ;  
 $X_5$  est  $-X_{10}$ ,  $-OX_{10}$ ,  $-SX_{10}$ ,  $-NX_8X_{10}$  ou  $-SO_2X_{11}$  ;  
 $X_6$  est hydrocarbyle substitué, hétéroaryle, un groupe protecteur d'hydroxy ou un groupe fonctionnel qui augmente la solubilité dans l'eau du dérivé de taxane ;  
 $X_7$  est hydrocarbyle, hydrocarbyle substitué, hétéroaryle ou un groupe protecteur de sulfhydryle ;  
 $X_8$  est l'hydrogène, hydrocarbyle ou hydrocarbyle substitué ;  
 $X_9$  est un groupe protecteur d'amino ;  
 $X_{10}$  est hydrocarbyle, hydrocarbyle substitué ou hétéroaryle ;  
 $X_{11}$  est hydrocarbyle, hydrocarbyle substitué, hétéroaryle,  $-OX_{10}$  ou  $-NX_8X_{14}$  ; et  
 $X_{14}$  est l'hydrogène, hydrocarbyle, hydrocarbyle substitué ou hétéroaryle.

5. Procédé selon la 4 où  
 $R_1$  est hydroxy ou forme un carbonate avec  $R_{14}$  ou  $R_2$  ;  
 $R_2$  est  $-OCOZ_2$ ,  $-OCOOZ_2$  ou forme un carbonate avec  $R_1$  ;  
 $R_4$  est  $-OCOZ_4$  ;  
 $R_9$  est l'hydrogène ou céto ;  
 $R_{13}$  est hydroxy, hydroxy ou



- $R_{14}$  est l'hydrogène, hydroxy, hydroxy protégé ou forme un carbonate avec  $R_1$  ;  
 $X_1$  est  $-OX_6$  ou  $-NX_8X_9$  ;  
 $X_2$  est l'hydrogène, hydrocarbyle ou hydrocarbyle substitué ;  
 $X_3$  et  $X_4$  sont indépendamment l'hydrogène, hydrocarbyle, hydrocarbyle substitué ou hétéroaryle ;  
 $X_5$  est  $-X_{10}$ ,  $-OX_{10}$  ou  $-NX_8X_{10}$  ;  
 $X_6$  est un groupe protecteur d'hydroxy ;  
 $X_8$  est l'hydrogène, hydrocarbyle ou hydrocarbyle substitué ;  
 $X_9$  est un groupe protecteur d'amino ;  
 $X_{10}$  est hydrocarbyle, hydrocarbyle substitué ou hétéroaryle ; et  
 $Z_2$  et  $Z_4$  sont indépendamment hydrocarbyle, hydrocarbyle substitué ou hétéroaryle.

6. Procédé selon l'une quelconque des revendications 1 à 5 où le mélange réactionnel contient un acide de Lewis.
7. Procédé selon la revendication 6 où l'acide de Lewis est choisi parmi les halogénures ou triflates des éléments du groupe IB, IIB, IIIB, IVB, VB, VIIB, VIII, IIIA, IVA, des lanthanides et des actinides.
8. Procédé selon la revendication 7 où l'acide de Lewis est choisi parmi chlorure de zinc, le chlorure stannique, le

trichlorure de cérium, le chlorure cuivreux, le trichlorure de lanthane, le trichlorure dysprosium et le trichlorure d'ytterbium.

- 5 9. Procédé selon l'une quelconque des revendications 1 à 8 où le taxane en C(10) comprend un groupe hydroxy en C(7) et le procédé comprend en outre le traitement du taxane acylé en C(10) avec un agent silylant pour silyler le groupe hydroxy en C(7).
- 10 10. Procédé selon l'une quelconque des revendications 1 à 8 où le taxane en C(10) comprend un groupe hydroxy en et le procédé comprend en outre le traitement du taxane acylé en C(10) avec un agent acylant pour acyler le hydroxy en C(7).
11. Procédé selon la revendication 9 ou la revendication 10 où le taxane en C(10) est la baccatine III.
- 15 12. Procédé selon l'une des revendications 1 à 8 où le taxane acylé en C(10) un substituant hydroxy, oxyde métallique ou oxyde en C(13) et le procédé en outre l'étape d'estérification du taxane acylé en C(10) par du taxane acylé en C(10) avec un de chaîne latérale choisi parmi les  $\beta$ -lactames, les oxazolines, les acides oxazolidine carboxyliques, les anhydrides d'acides oxazolidine carboxyliques et les dérivés d'isosérine.

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